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enue, Rockville, MD 20851 (US). OLSEN, Henrik, S. [DKUIS]; 182 Kendrick Place #24, Galthersburg, MD 20878 (US). EBNER, Reinhard (DE/US); 9906 Shelburn Eromee #316, Galthersburg, MD 20878 (US). BREWER, Laurie, A. [US/US); 14920 Mount Nebo Road, Poolesville, MD 20837 (US). MOORE, Paul, A. [GB/US]; Apartment #104, 1908 Holly Ridge Drive, McLean, VA 22102 (US). SHI, Yanggu [CN/US]; 437 West Sido Drive, Galthersburg, MD 20878 (US), LAFLEUR, David, W. [US/US]; 1615 Q Street, N. W. #807, Washington, DC 20009 (US). LI, Y1 [CN/US]; 1247 Lakeide Drive, #3034, Sumnyrale, CA 94086 (US). ZENG, MD 20878 (US); 1950 Saddleview Drive, Galthersburg, MD 20878 (US); XYAW, Hia [BU/US]; 220 Sugarbush Cirke, Perderick, MD 2070 (US). Drive, Bethesda, MD 20814 (US). HÜ, Jing-Shan [CN/US]: 1247 Lakeside Drive #3034, Sunnyvale, CA 94086 (US). FLORENCE, Kimberly, A. [US/US]: 12805 Atlantic Avcle, Frederick, MD 21703 (US). MD 20874 (US). DUAN, Roxanne [US/US]; 4541 Fairfield

(74) Agents: BROOKES, Anders; A. et al.; Human Genome Sciences, Inc., 9410 Key West Avenue, Rockville, MD 10850 (US).

(71) Applicant (for all designated States except US): HUMAN GENOME SCIENCES, INC. [US/US]; 9410 Key West Avenue, Rockville, MD 20850 (US).

(Continued on the following page)

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(72) Inventors; and
(75) Inventors; and
(75) Inventors/Applicants (for US only): RUBEN, Steven, M. (USUS); 18528 Heritage Hills Drive, Olney, MD 20832 (US), ROSEN, Craig, A. (USUS); 22400 Rolling Hills Road, Laytonsville, MD 20882 (US), FISCHER, Carrie, L. (USUS); 8301 Hall Street, Burke, VA 22015 (US), SOP-PET, Daniel, R. (USUS); 15050 Stillfield, Place, Centreville, VA 22020 (US), CARTER, Kenneth, C. (USUUS); 11601 Brandy Hall Lane, North Potomac, MD 20878 (US), BEDNARIK, Daniel, P. (USUUS); 8822 Blue Sea Drive, Columbia, MD 21046 (US). RUREES, Gregory, A. (USUUS); 9729 Clagett Farm Drive, Potomac, MD 20854 (US), PU, Guo-Liang (CN/US); 13524 Straw Bale Lane, Damestown, MD 20878 (US), NJ, Jian (CN/US); 5002 Manorfield Road, Rockville, MD 20853 (US), FENG, Ping (CN/US); 4 Reida Court, Gaithersburg, MD 20878 (US), YOUNG, Paul, E. (US/US); 122 Beckwith Street, Gaithersburg, MS 20878 (US). 

Published

burg, MD 20878 (US). GREENE, John, M. (US/US); 872 Diamond Drive, Galthersburg, MD 20878 (US). FERRIE, Ann, M. [US/US]; 13203 L Astoria Hill Court, Germantown,

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(54) Title: 186 HUMAN SECRETED PROTEINS

(57) Abstract

The present invention relates to 186 novel human secreted proteins and Isolated nucleic acids containing the coding regions of the genes encoding such proteins. Also provided are vectors, host cells, antibodies, and recombinant methods for producing human secreted proteins. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to these novel human secreted proteins.

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#### 186 Human Secreted Proteins

#### Field of the Invention

This invention relates to newly identified polynucleotides and the polypeptides encoded by these polynucleotides, uses of such polynucleotides and polypeptides, and their production.

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#### Background of the Invention

Unlike bacterium, which exist as a single compartment surrounded by a membrane, human cells and other eucaryotes are subdivided by membranes into many functionally distinct compartments. Each membrane-bounded compartment, or organelle, contains different proteins essential for the function of the organelle. The cell uses "sorting signals," which are amino acid motifs located within the protein, to target proteins to particular cellular organelles.

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One type of sorting signal, called a signal sequence, a signal peptide, or a leader sequence, directs a class of proteins to an organelle called the endoplasmic reticulum (ER). The ER separates the membrane-bounded proteins from all other types of proteins. Once localized to the ER, both groups of proteins can be further directed to another organelle called the Golgi apparatus. Here, the Golgi distributes the proteins to vesicles, including secretory vesicles, the cell membrane, lysosomes, and the other organelles.

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Proteins targeted to the ER by a signal sequence can be released into the extracellular space as a secreted protein. For example, vesicles containing secreted proteins can fuse with the cell membrane and release their contents into the extracellular space - a process called exocytosis. Exocytosis can occur constitutively or after receipt of a triggering signal. In the latter case, the proteins are stored in secretory vesicles (or secretory granules) until exocytosis is triggered. Similarly, proteins residing on the cell membrane can also be secreted into the extracellular space by proteolytic cleavage of a "linker" holding the protein to the membrane.

Despite the great progress made in recent years, only a small number of genes encoding human secreted proteins have been identified. These secreted proteins include the commercially valuable human insulin, interferon, Factor VIII, human growth hormone, tissue plasminogen activator, and erythropoeitin. Thus, in light of the pervasive role of secreted proteins in human physiology, a need exists for identifying and characterizing novel human secreted proteins and the genes that encode them. This knowledge will allow one to detect, to treat, and to prevent medical disorders by using secreted proteins or the genes that encode them.

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#### Summary of the Invention

The present invention relates to novel polynucleotides and the encoded polypeptides. Moreover, the present invention relates to vectors, host cells, antibodies, and recombinant methods for producing the polypeptides and polynucleotides. Also provided are diagnostic methods for detecting disorders related to the polypeptides, and therapeutic methods for treating such disorders. The invention further relates to screening methods for identifying binding partners of the polypeptides.

#### Detailed Description

#### Definitions

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The following definitions are provided to facilitate understanding of certain terms used throughout this specification.

In the present invention, "isolated" refers to material removed from its original environment (e.g., the natural environment if it is naturally occurring), and thus is altered "by the hand of man" from its natural state. For example, an isolated polynucleotide could be part of a vector or a composition of matter, or could be contained within a cell, and still be "isolated" because that vector, composition of matter, or particular cell is not the original environment of the polynucleotide.

In the present invention, a "secreted" protein refers to those proteins capable of being directed to the ER, secretory vesicles, or the extracellular space as a result of a signal sequence, as well as those proteins released into the extracellular space without necessarily containing a signal sequence. If the secreted protein is released into the extracellular space, the secreted protein can undergo extracellular processing to produce a "mature" protein. Release into the extracellular space can occur by many mechanisms, including exocytosis and proteolytic cleavage.

As used herein, a "polynucleotide" refers to a molecule having a nucleic acid sequence contained in SEQ ID NO:X or the cDNA contained within the clone deposited with the ATCC. For example, the polynucleotide can contain the nucleotide sequence of the full length cDNA sequence, including the 5' and 3' untranslated sequences, the coding region, with or without the signal sequence, the secreted protein coding region, as well as fragments, epitopes, domains, and variants of the nucleic acid sequence. Moreover, as used herein, a "polypeptide" refers to a molecule having the translated amino acid sequence generated from the polynucleotide as broadly defined.

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In the present invention, the full length sequence identified as SEQ ID NO.X was often generated by overlapping sequences contained in multiple clones (contig

analysis). A representative clone containing all or most of the sequence for SEQ ID NO:X was deposited with the American Type Culture Collection ("ATCC"). As shown in Table 1, each clone is identified by a cDNA Clone ID (Identifier) and the ATCC Deposit Number. The ATCC is located at 12301 Park Lawn Drive, Rockville Maryland 20852, USA. The ATCC deposit was made pursuant to the terms of the

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purposes of patent procedure.

Budapest Treaty on the international recognition of the deposit of microorganisms for

A "polynucleotide" of the present invention also includes those polynucleotides capable of hybridizing, under stringent hybridization conditions, to sequences contained in SEQ ID NO:X, the complement thereof, or the cDNA contained within the clone deposited with the ATCC. "Stringent hybridization conditions" refers to an overnight incubation at 42° C in a solution comprising 50% formamide, 5x SSC (750 mM NaCl, 75 mM sodium citrate), 50 mM sodium phosphate (pH 7.6), 5x Denhardt's solution, 10% dextran sulfate, and 20 µg/ml denatured, sheared salmon sperm DNA, followed

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15 by washing the filters in 0.1x SSC at about 65°C.
Also contemplated are nucleic acid molecules that hybridize to the polynucleotides of the present invention at lower stringency hybridization conditions. Changes in the stringency of hybridization and signal detection are primarily

of formamide result in lowered stringency); salt conditions, or temperature. For example, lower stringency conditions include an overnight incubation at 37°C in a solution comprising 6X SSPE (20X SSPE = 3M NaCl; 0.2M NaH,PO,; 0.02M EDTA pH 7.4), 0.5% SDS, 30% formamide, 100 ug/ml salmon sperm blocking DNA;

accomplished through the manipulation of formamide concentration (lower percentages

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followed by washes at 50°C with 1XSSPE, 0.1% SDS. In addition, to achieve even

25 lower stringency, washes performed following stringent hybridization can be done at higher salt concentrations (e.g. 5X SSC).

Note that variations in the above conditions may be accomplished through the inclusion and/or substitution of alternate blocking reagents used to suppress background in hybridization experiments. Typical blocking reagents include Denhardt's reagent, BLOTTO, heparin, denatured salmon sperm DNA, and commercially available proprietary formulations. The inclusion of specific blocking reagents may require modification of the hybridization conditions described above, due to problems with compatibility.

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Of course, a polynucleotide which hybridizes only to polyA+ sequences (such as any 3' terminal polyA+ tract of a cDNA shown in the sequence listing), or to a

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complementary stretch of T (or U) residues, would not be included in the definition of "polynucleotide," since such a polynucleotide would hybridize to any nucleic acid molecule containing a poly (A) stretch or the complement thereof (e.g., practically any double-stranded cDNA clone).

5 5 S or for other reasons. "Modified" bases include, for example, tritylated bases and double-stranded regions, hybrid molecules comprising DNA and RNA that may be and double-stranded DNA, DNA that is a mixture of single- and double-stranded or modified RNA or DNA. For example, polynucleotides can be composed of singlepolyribonucleotide or polydeoxribonucleotide, which may be unmodified RNA or DNA RNA; thus, "polynucleotide" embraces chemically, enzymatically, or metabolically unusual bases such as inosine. A variety of modifications can be made to DNA and contain one or more modified bases or DNA or RNA backbones modified for stability regions comprising RNA or DNA or both RNA and DNA. A polynucleotide may also stranded regions. In addition, the polynucleotide can be composed of triple-stranded single-stranded or, more typically, double-stranded or a mixture of single- and doubleregions, single- and double-stranded RNA, and RNA that is mixture of single- and modified forms. The polynucleotide of the present invention can be composed of any

ટ્ટ 20 ઝ ဗ as well as in a voluminous research literature. Modifications can occur anywhere in a to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and branched, for example, as a result of ubiquitination, and they may be cyclic, with or polypeptide, including the peptide backbone, the amino acid side-chains and the amino Such modifications are well described in basic texts and in more detailed monographs polypeptides may be modified by either natural processes, such as posttranslational may contain amino acids other than the 20 gene-encoded amino acids. The nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, given polypeptide may contain many types of modifications. Polypeptides may be or carboxyl termini. It will be appreciated that the same type of modification may be processing, or by chemical modification techniques which are well known in the art present in the same or varying degrees at several sites in a given polypeptide. Also, a formation, demethylation, formation of covalent cross-links, formation of cysteine, covalent attachment of phosphotidylinositol, cross-linking, cyclization, disulfide bond attachment of flavin, covalent attachment of a heme moiety, covalent attachment of a Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent from posttranslation natural processes or may be made by synthetic methods. without branching. Cyclic, branched, and branched cyclic polypeptides may result The polypeptide of the present invention can be composed of amino acids joined

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formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination, methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, transfer-RNA mediated addition of amino acids to proteins such as arginylation, and ubiquitination. (See, for instance, PROTEINS - STRUCTURE AND MOLECULAR PROPERTIES, 2nd Ed., T. E. Creighton, W. H. Freeman and Company, New York (1993); POSTTRANSLATIONAL COVALENT MODIFICATION OF PROTEINS, B. C. Johnson, Ed., Academic Press, New York, pgs. 1-12 (1983); Seifter et al., Meth Enzymol 182:626-646 (1990);

"SEQ ID NO:X" refers to a polynucleotide sequence while "SEQ ID NO:Y" refers to a polypeptide sequence, both sequences identified by an integer specified in Table 1.

Rattan et al., Ann NY Acad Sci 663:48-62 (1992).)

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"A polypeptide having biological activity" refers to polypeptides exhibiting activity similar, but not necessarily identical to, an activity of a polypeptide of the present invention, including mature forms, as measured in a particular biological assay, with or without dose dependency. In the case where dose dependency does exist, it need not be identical to that of the polypeptide, but rather substantially similar to the dose-dependence in a given activity as compared to the polypeptide of the present invention (i.e., the candidate polypeptide will exhibit greater activity or not more than about 25-fold less and, preferably, not more than about tenfold less activity, and most preferably, not more than about three-fold less activity relative to the polypeptide of the present invention.)

#### 25 Polynucleotides and Polypeptides of the Invention

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 1

This gene is expressed primarily in testes tumor and to a lesser extent in fetal brain.

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer particularly of the testes, and defects of the central nervous system such as seizure and neurodegenerative disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues of cells, particularly cancer of the testes and central nervous system,

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expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., testes and other reproductive tissue, brain and other tissue of the nervous system, and blood cells, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of testicular cancer and treatment of central nervous system disorders since this gene is primarily expressed in the testes tumor and developing brain.

or bodily fluid from an individual not having the disorder.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 2

This gene is expressed primarily in cancer tissues, such as breast cancer and Wilm's tumor, and to a lesser extent in fetal tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, and/or tumors, particularly, those found in the breast, and developmental

- abnormalities or disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the glandular tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., mammary tissue, and fetal tissue and, cancerous and wounded tissues) or bodily fluids (e.g.,
  - serum, plasma, urine, synovial fluid or spinal fluid) or another tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 314 as residues: Pro-11 to Thr-18, Leu-43 to Pro-50,
- The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of cancers and/or tumors, particularly, those found in the breast since expression is mainly in cancer/tumor tissues. May serve as therapeutic proteins for proliferation/differentiation of letal

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Gly-64 to Leu-72, and Leu-81 to Lys-86.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 3

extent in spleen, chronic lymphocytic leukemia. This gene is expressed primarily in CD34 depleted buffy coat and to a lesser

5 5 S significantly higher or lower levels may be routinely detected in certain tissues and cell differential identification of the tissue(s) or cell type(s). For a number of disorders of biological sample and for diagnosis of diseases and conditions: blood disorders or cell sample taken from an individual having such a disorder, relative to the standard the above tissues or cells, particularly of the immune system, expression of this gene at directed to these polypeptides are useful in providing immunological probes for leukemias, diseases of the immune system. Similarly, polypeptides and antibodies reagents for differential identification of the tissue(s) or cell type(s) present in a gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an types (e.g., blood cells, and spleen, and cancerous and wounded tissues) or bodily individual not having the disorder. fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or Therefore, polynucleotides and polypeptides of the invention are useful as

corresponding to this gene are useful for treatment/diagnosis of blood disorders or leukemias, diseases of the immune system since expression is in tissues related to The tissue distribution indicates that polynucleotides and polypeptides

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 4

This gene is expressed primarily in CD34 depleted buffy coal

ઝ 25 ઝ of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells biological sample and for diagnosis of diseases and conditions: blood disorders or reagents for differential identification of the tissue(s) or cell type(s) present in a expression level in healthy tissue or bodily sluid from an individual not having the having such a disorder, relative to the standard gene expression level, i.e., the synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, lower levels may be routinely detected in certain tissues and cell types (e.g., blood particularly of the immune system, expression of this gene at significantly higher or polypeptides are useful in providing immunological probes for differential identification lymphocytic diseases. Similarly, polypeptides and antibodies directed to these Therefore, polynucleotides and polypeptides of the invention are useful as

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expression is in tissues related to immune function. corresponding to this gene are useful for treatment/diagnosis of blood disorders since The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 5

This gene is expressed primarily in CD34 depleted buffy coat

20 5 5 and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or or cell type(s). For a number of disorders of the above tissues or cells, particularly of diseases. Similarly, polypeptides and antibodies directed to these polypeptides are Pro-13 to Lys-21. epitopes include those comprising a sequence shown in SEQ ID NO. 317 as residues: spinal fluid) or another tissue or cell sample taken from an individual having such a be routinely detected in certain tissues and cell types (e.g., blood cells, and cancerous the immune system, expression of this gene at significantly higher or lower levels may useful in providing immunological probes for differential identification of the tissue(s) biological sample and for diagnosis of diseases and conditions: blood or immune reagents for differential identification of the tissue(s) or cell type(s) present in a healthy tissue or bodily fluid from an individual not having the disorder. Preferred disorder, relative to the standard gene expression level, i.e., the expression level in Therefore, polynucleotides and polypeptides of the invention are useful as

expression is in tissues related to immune function. corresponding to this gene are useful for treatment/diagnosis of blood disorders since The tissue distribution indicates that polynucleotides and polypeptides

#### 23 FEATURES OF PROTEIN ENCODED BY GENE NO: 6

This gene is expressed primarily in CD34 depleted buffy coat

diseases. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: blood or immune reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

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- cancerous and wounded tissues) or bodily sluids (e.g., serum, plasma, urine, synovial or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) be routinely detected in certain tissues and cell types (e.g., and blood cells, and the immune system, expression of this gene at significantly higher or lower levels may fluid or spinal fluid) or another tissue or cell sample taken from an individual having
- ઝ such a disorder, relative to the standard gene expression level, i.e., the expression level

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in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 318 as residues: Lys-31 to Lys-39.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment/diagnosis of blood diseases since it is expressed in tissues related to immune function.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 7

This gene is expressed primarily in CD34 depleted buffy coat and to a lesser extent in pineal gland.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases of the immune system and brain associated diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and pineal gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another

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standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatmen/diagnosis of blood disorders, immune diseases or brain associated diseases (specifically of the pineal gland) since

issue or cell sample taken from an individual having such a disorder, relative to the

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 8

expression is in tissues related to immune function.

The translation product of this gene shares sequence homology with an organic cation transporter which is thought to be important in organic cation uptake in the kidney and liver. (See Accession No. 2343059.) Preferred polypeptide fragments comprise the amino acid sequence ITIAIQMICLYNXELYPTFVRNXGVMVCSSLCDIGGIITP FIVFRLREVWQALPLILFAVLGLLAGGYTLLPETKGVALPETMKDAENLGRKAKPKENTYLK 35 VQTSEPSGT (SEQ ID NO: 615) or TMKDAENLGRKAKPKENT (SEQ ID NO: 616) as well as N-terminal and C-terminal deletions of these fragments. Also preferred are polynucleotide fragments encoding these polypeptide fragments.

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This gene is expressed primarily in liver.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hepatic and renal diseases where drug elimination/cation exchange (organic cation uptake) in the liver and kidney are problematic. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hepatic or renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., kidney

10 or lower levels may be routinely detected in certain tissues and cell types (e.g., kidney and liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 320 as residues: Asn-64 to Asn-74, and Gln-81 to Gly-87.

The tissue distribution and homology to organic cation transporter indicate that polynucleotides and polypeptides corresponding to this gene are useful as a polyspecific transporter that is important for drug elimination in the liver (and possibly kidney) since expression is found in the liver.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 9

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This gene is expressed primarily in eosinophil induced with IL-5 and to a lesser extent in fetal liver and spleen. This gene also maps to chromosome 15, and therefore can be used in linkage analysis as a marker for chromosome 15.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases of the immune system, particularly allergies or asthma. Similarly, polypeptides and antibodies directed

to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, liver, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the

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standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

esosinphil reactions since expression seems to be concentrated in eosinophils and other corresponding to this gene are useful for treating/diagnosis of diseases involving The tissue distribution indicates that polynucleotides and polypeptides

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 10

5 lesser extent in Hodgkins lymphoma. Any frame shifts in this sequence can easily be clarified using known molecular biology techniques. This gene is expressed primarily in tissues of hematopoietic lineage and to a

biological sample and for diagnosis of diseases and conditions which include, but are reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

- 20 5 cancerous tissues) or bodily sluids (e.g., serum, plasma, urine, synovial fluid or spinal cell types (e.g., hematopoietic cells, lymphoid and reticuloendothelial tissues, and at significantly higher or lower levels may be routinely detected in certain tissues and of the above tissues or cells, particularly of the immune system, expression of this gene antibodies directed to these polypeptides are useful in providing immunological probes relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, for differential identification of the tissue(s) or cell type(s). For a number of disorders
- 23 corresponding to this gene are useful for treatment diagnosis for lymphomas or expression in anergic T-cells and lymphomas. immune dysfuction or as a therapeutic protein useful in immune modulation based on The tissue distribution indicates that polynucleotides and polypeptides

#### 30 FEATURES OF PROTEIN ENCODED BY GENE NO: 11

This gene is expressed primarily in neutrophils and to a lesser extent in activated

for diagnosis of diseases and conditions: inflamation. Similarly, polypeptides and reagents for differential identification of the cell type present in a biological sample and Therefore, polynucleotides and polypeptides of the invention are useful as

for differential identification of the tissue(s) or cell type(s). For a number of disorders antibodies directed to these polypeptides are useful in providing immunological probes ઝ

not limited to, and immune deficiency or dysfunction. Similarly, polypeptides and tissues involved in immunity such as the liver and spleen. or bodily fluid from an individual not having the disorder. PCT/US98/04493 25 8 2 8 5

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or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another at significantly higher or lower levels may be routinely detected in certain tissues and comprising a sequence shown in SEQ ID NO. 323 as residues: Glu-40 to Lys-46. standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the cell types (e.g., blood cells and lymphoid tissue, and cancerous and wounded tissues) of the above tissues or cells, particularly of the immune system, expression of this gene fluid from an individual not having the disorder. Preferred epitopes include those

corresponding to this gene are useful for modulation of an immune reaction or as a growth factor for the differentiation or proliferation of neutrphils for the treatment of

The tissue distribution indicates that polynucleotides and polypeptides

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 12

continues in the 5' direction. Preserred polypeptide fragments comprise the amino acid sequence PRVRNSPEDLGLSLTGDSCKL (SEQ ID NO:617). cells. It is likely that the open reading frame containing the predicted signal peptide This gene is expressed primarily in brain and to a lesser extent in activated T-

number of disorders of the above tissues or cells, particularly of the central nervous reagents for differential identification of the tissue(s) or cell type(s) present in a an individual having such a disorder, relative to the standard gene expression level, i.e., plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from detected in certain tissues and cell types (e.g., blood cells, and brain, and other tissue of system, expression of this gene at significantly higher or lower levels may be routinely disorders including ischemic shock, alzheimers and cognitive disorders. Similarly Val-96, Lys-136 to Ser-145, Ile-152 to Met-169, and Arg-189 to Lys-196 NO. 324 as residues: Ser-5 to Glu-14, Ile-21 to Pro-35, Ser-65 to Asp-81, Cys-89 to disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID the expression level in healthy tissue or bodily fluid from an individual not having the the nervous system and cancerous and wounded tissues) or bodily fluids (e.g., serum, immunological probes for differential identification of the tissue(s) or cell type(s). For a polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: neurodegenerative Therefore, polynucleotides and polypeptides of the invention are useful as

tissue or in the treatment of neurological disorders of the CNS. corresponding to this gene are useful for diagnostic/treatment for cancers of the given The tissue distribution indicates that polynucleotides and polypeptides

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 13

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This gene was also recently cloned by other groups, naming this calcium-activated potassium channel gene, hKCa4. (See Accession No. AF033021, see also, Accession No. 2584866.) This gene is mapped to human chromosome 19q13.2. A second signal sequence likely exists upstream from the predicted signal sequence as described in Table I. Preferred polypeptide fragments comprise: QADDLQATVAALCVLRGGGPWAG SWLSPKTPGAMGGDLVLGGALRRRKRLL (SEQ NO: 618); or EQEKSLAGWALVLAXXGIGL MVLHAEMLWFGGCSAVNATGHLSDTLWLIPTFLTIGYGDVVPCTMWGKIVCLCTGVMGVCC TALLVAVVARKLEFNKAEKHVHNFMMDIQYTKEMKESAARVLQEAWMFYKHTRRKESHAAR XHQRXLLAAINAFRQVRLKHRKLREQVNSMVDISKMHMILYDLQQNLSSSHRALEKQIDTLAG KLDALTELLSTALGPRQLPEPSQQSK (SEQ ID NO: 619), as well as N-terminal and C-terminal deletions. Also preferred are polynulcleotide fragments encoding these polypeptide fragments.

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15 This gene is expressed primarily in breast lymph node and T-cells, and to a lesser extent in placenta.

immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides lower levels may be routinely detected in certain tissues and cell types (e.g., lymphoid issue, blood cells and placenta, and cancerous and wounded tissues) or bodily fluids sample taken from an individual having such a disorder, relative to the standard gene particularly of the immune system, expression of this gene at significantly higher or expression level, i.e., the expression level in healthy tissue or bodily fluid from an (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell Therefore, polynucleotides and polypeptides of the invention are useful as ndividual not having the disorder. Preferred epitopes include those comprising a are useful in providing immunological probes for differential identification of the biological sample and for diagnosis of diseases and conditions: hematologic and issue(s) or cell type(s). For a number of disorders of the above tissues or cells, reagents for differential identification of the tissue(s) or cell type(s) present in a sequence shown in SEQ ID NO. 325 as residues: Arg-13 to Lys-23. 52 2 3

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment/diagnosis of hematologic and diseases involving immune modulation based or distribution in the lymph node and T-cells.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 14

This gene was recently cloned by another group, calling it PAPS synethase. (See Accession No. e1204135.) Preferred polypeptide fragments comprise the amino acid sequence YQAHHVSRNKRGQVVGTRGGFRGCTVWLTGLSGAGK (SEQ ID NO: 620).

Also preferred are the polynucleotide fragments encoding this polypeptide fragment.

It has been discovered that this gene is expressed primarily in benign prostate hyperplasia, Human Umbilical Vein Endothelial Cells and to a lesser extent in smooth muscle and Human endometrial stromal cells-treated with estradiol.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: inflamation, ischemia, and restenosis, based on endothelial cell and smooth muscle cell expression, and prostate diseases such as benign prostate hyperplasia or prostate cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a

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immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate or vessels of the circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., prostate, endothelial cells, smooth muscle, and endometrium, and cancerous and wounded tissues) or bodily

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fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO. 326 as residues: Arg-21 to Asp-26, Lys-35 to Lys-44, Glu-49 to Asn-58.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating/diagnosing diseases or conditions where the endothelial cell lining of the veins and arteries of underlying smooth muscle are involved.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 15

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This gene is expressed primarily in human 6 week embryo and to a lesser extent placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: developmental anomalies or fetal deficiencies. Similarly, polypeptides and antibodies directed to these

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NO. 327 as residues Lys-50 to Glu-57. an individual having such a disorder, relative to the standard gene expression level, i.e., disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from tissue, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, the expression level in healthy tissue or bodily fluid from an individual not having the particularly developmental in nature, expression of this gene at significantly higher or polypeptides are useful in providing immunological probes for differential identification lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic

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corresponding to this gene are useful for detection of developmental abnormalities. The tissue distribution indicates that polynucleotides and polypeptides 5

#### FEATURES OF PROTEIN ENCODED BY GENE NO:

2 linkage analysis as a marker for chromosome 14. in fetal tissues. This gene is mapped to chromosome 14, and therefore is useful in This gene is expressed primarily in kidney and amygdala and to a lesser extent

25 20 system or developing fetal tissues, expression of this gene at significantly higher or of the tissue(s). For a number of disorders of the above tissues, particularly of the renal polypeptides are useful in providing immunological probes for differential identification developmental abnormalities. Similarly, polypeptides and antibodies directed to these for diagnosis of diseases and conditions: kidney diseases, neurological disorders and reagents for differential identification of the tissue(s) present in a biological sample and Therefore, polynucleotides and polypeptides of the invention are useful as

30 serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample arnygdala, and fetal tissues, and cancerous and wounded tissues) or bodily fluids (e.g., lower levels may be routinely detected in certain tissues and cell types (e.g., kidney, individual not having the disorder. expression level, i.e., the expression level in healthy tissue or bodily fluid from an taken from an individual having such a disorder, relative to the standard gene

corresponding to this gene are useful for treatment or diagnosis of conditions affecting the brain, kidneys and fetal development The tissue distribution indicates that polynucleotides and polypeptides

#### ၓ FEATURES OF PROTEIN ENCODED BY GENE NO: 17

This gene is expressed primarily in ovarian cancer

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5 plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) ovarian cancer Similarly, polypeptides and antibodies directed to these polypeptides are the expression level in healthy tissue or bodily fluid from an individual not having the an individual having such a disorder, relative to the standard gene expression level, i.e. reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, may be routinely detected in certain tissues and cell types (e.g., ovarian and other the reproductive system. expression of this gene at significantly higher or lower levels biological sample and for diagnosis of diseases and conditions; solid tumors similar to reagents for differential identification of the tissue(s) or cell type(s) present in a NO. 329 as residues Ser-51 to Val-56. disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID Therefore, polynucleotides and polypeptides of the invention are useful as

corresponding to this gene are useful for the treatment of solid tumors of the reproductive system such as ovarian cancer. The tissue distribution indicates that polynucleotides and polypeptides

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 18

polypeptide fragments. and C-terminal deletions. Also preferred are polynucleotide fragments encoding these GLTLTTCSGPTEKPATKNYFLKRLLQEMHIRAN (SEQ ID NO: 644), as well as N-terminal ILILPYCAHLHEELNC (SEQ ID NO: 643) and SFFISEEKGHLLLQAERHPWVAGALVGVSG fragments comprise the amino acid sequence: IRHEQHPNFSLEMHSKGSSLLLFLPQL This gene is expressed primarily in brain medulloblastoma, Preferred polypeptide

25 20 30 or cell type(s). For a number of disorders of the above tissues or cells, particularly of biological sample and for diagnosis of diseases and conditions: tumors particularly of sample taken from an individual having such a disorder, relative to the standard gene tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids levels may be routinely detected in certain tissues and cell types (e.g., brain and other the Central nervous system, expression of this gene at significantly higher or lower useful in providing immunological probes for differential identification of the tissue(s) the CNS or Similarly, polypeptides and antibodies directed to these polypeptides are reagents for differential identification of the tissue(s) or cell type(s) present in a (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell Therefore, polynucleotides and polypeptides of the invention are useful as

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expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating medulloblastoma or similar tumors.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 19

This gene is expressed primarily in adipocytes.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: obesity. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the adipose tissues expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., adipocytes and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating obesity by regulating the function and number of adipocytes

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 20

This gene is expressed primarily in B cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, of the immune system with an emphasis on B cell lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the tumors of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of B cell derived

5 tumors based on its expression in b cell lymphomas

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 21

This gene is expressed primarily in immune cells and to a lesser extent in fetal tissues

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: inflammatory diseases Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., cells of the immune system; and fetal tissues, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an

individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. Preferred epitopes include those comprising a sequence shown in SEQ ID NO:333 as residues Asp-10 to Pro-19, Ser-74 to Tyr-79, Glu-95 to Lys-110.

The tissue distribution indicates that polynucleotides and polypeptides

corresponding to this gene are useful for treatment of diseases involving alterations in T

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 22

It has been discovered that this gene is expressed primarily in ovarian tumor. Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors particularly of the ovary. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of

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tumors of the reproductive organs. expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., ovarian

sequence shown in SEQ ID NO. 334 as residues: Leu-22 to Gln-27 individual not having the disorder. Preferred epitopes include those comprising a expression level, i.e., the expression level in healthy tissue or bodily fluid from an sample taken from an individual having such a disorder, relative to the standard gene and other reproductive tissue and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell

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has only been identified in ovarian tumors corresponding to this gene are useful for diagnosis and treatment of ovarian tumors as it The tissue distribution indicates that polynucleotides and polypeptides

### FEATURES OF PROTEIN ENCODED BY GENE NO: 23

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a lesser extent in osteoclastoma cell line It has been discovered that this gene is expressed primarily in fetal tissues and to

2 Similarly, polypeptides and antibodies directed to these polypeptides are useful in biological sample and for diagnosis of diseases and conditions: osteoporosis or arthritis reagents for differential identification of the tissue(s) or cell type(s) present in a providing immunological probes for differential identification of the tissue(s) or cel type(s). For a number of disorders of the above tissues or cells, particularly of the Therefore, polynucleotides and polypeptides of the invention are useful

skeletal expression of this gene at significantly higher or lower levels may be routinely in healthy tissue or bodily fluid from an individual not having the disorder. cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial detected in certain tissues and cell types (e.g., bone cells, and fetal tissue, and such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having

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corresponding to this gene are useful for treatment of conditions of abnormal bone marker for malignancies derived from osteoclasts or their precursors. remodeling due to enhanced activity of osteoclasts. This may be useful as a specific The tissue distribution indicates that polynucleotides and polypeptides 23

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 24

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polynucleotides periplusmic ribonuclease which is thought to be important in degrading extracellular The translation product of this gene shares sequence homology with a

smooth muscle cells It has been discovered that this gene is expressed primarily in serum treated

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5 S spinal fluid) or another tissue or cell sample taken from an individual having such a the vasculature expression of this gene at significantly higher or lower levels may be or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) restenosis. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: vascular disease such as reagents for differential identification of the tissue(s) or cell type(s) present in a healthy tissue or bodily fluid from an individual not having the disorder. Preferred routinely detected in certain tissues and cell types (e.g., smooth muscle, and cancerous Gln-30 to Lys-36, and Pro-41 to Arg-48. epitopes include those comprising a sequence shown in SEQ ID NO: 336 as residues: disorder, relative to the standard gene expression level, i.e., the expression level in and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or Therefore, polynucleotides and polypeptides of the invention are useful as

2 polynucleotides and polypeptides corresponding to this gene are useful for treatment of pathological conditions of smooth muscle associated with bacterial or viral infiltration The tissue distribution and homology to ribonucleases indicate that

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 25

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မွ 25 cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial expression of this gene at significantly higher or lower levels may be routinely detected tissues or cells, particularly of the human brain development and related diseases, identification of the tissue(s) or cell type(s). For a number of disorders of the above these polypeptides are useful in providing immunological probes for differential development and related diseases. Similarly, polypeptides and antibodies directed to biological sample and for diagnosis of diseases and conditions: human brain reagents for differential identification of the tissue(s) or cell type(s) present in a such a disorder, relative to the standard gene expression level, i.e., the expression level in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and in healthy tissue or bodily fluid from an individual not having the disorder. fluid or spinal fluid) or another tissue or cell sample taken from an individual having This gene is expressed primarily in Early Stage Human Brain Therefore, polynucleotides and polypeptides of the invention are useful as

35 and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases affecting human brain development and related diseases. The tissue distribution and homology to this gene indicate that polynucleotides

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 26

It has been discovered that this gene is expressed primarily in human brain tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: human brain diseases and other diseases related to brain diseases, which may be caused by brain diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the human brain diseases, expression of this gene at significantly higher or lower levels

plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of

the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum,

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The tissue distribution and homology to the gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of human brain diseases and other diseases related.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 27

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune diseases, inflammatory diseases and diseases related to T lymph cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune diseases, inflammatory diseases and diseases related to T lymph cells, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample

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expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the gene indicate that polyhucleotides and polypeptides corresponding to this gene are useful for immune diseases,

5 inflammatory diseases and diseases related to Tlymph cells.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 28

The translation product of this gene shares sequence homology with Shigella flexneri positive transcriptional regulator CriR (criR) gene which is thought to be important in regulation of gene expression.

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This gene is expressed primarily in human synovial sarcoma and normal human brain tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: human brain diseases particularly sarcomas of the synovium. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the human brain and synovium and other related human brain diseases.

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brain diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain (e.g., synovial tissue, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of human synovial sarcoma and other related human brain diseases.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 29

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This gene is expressed in bone marrow, infant brain, fetal liver and spleen, prostate and to a lesser extent in pineal gland, adipose tissue, kidney, adrenal gland, umbilical vein endothelial cells, and T cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diseases related to bone marrow or

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taken from an individual having such a disorder, relative to the standard gene

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hematoplastic tissues, prostate, kidney, adrenal gland, and cardiovascular tissue or organs. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the diseases related to

- 5 hematoplastic tissues, immune system, prostate, kidney, adrenal gland, and cardiovascular tissue or organs, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, hematopoietic cells, pineal gland, adipose tissue, kidney, adrenal gland, endothelial cells, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an
- The tissue distribution and homology to the gene indicate that polynucleotides

  15 and polypeptides corresponding to this gene are useful for diagnosis and treatment of

  diseases related to hematoplastic tissues, immune system, prostate, kidney, adrenal

  gland, and cardiovascular tissue or organs.

individual not having the disorder.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 30

20 This gene is expressed primarily in meningea and to a lesser extent in breast and adult brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Diseases of the meanings and related brain diseases. Similarly polypeptides and antibodies directed

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meningea and related brain diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the meningea and related brain diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., miningea, mammary tissue, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the meningea and related brain diseases.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 3

This gene is expressed in meningea, fetal spleen, osteoblast and to a lesser extent in activated T-cells, endometrial stromal cells, fetal lung, HL-60, thymus, testis and endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: meningeal disease, osteoporosis, immune diseases, and hematoplastic diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell type(s). For a number of disorders of the

diseases, and hematoplastic diseases, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, endometrium, lung, thymus, testis, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of meningeal, osteoporosis, immune diseases, hematoplastic diseases, testis diseases and

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lung diseases.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 32

This gene is expressed primarily in human thymus and to a much lesser extent in infant brain, T-cells, smooth muscle, endothelial cells, bone marrow, human ovarian tumor and keratinocytes testes, osteoclastoma, breast, and tonsils.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Diseases involving the

biological sample and for diagnosis of diseases and conditions: Diseases involving the thymus, particularly thymic cancer and diseases involving T-cell maturation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a

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number of disorders of the above tissues or cells, particularly of the thymus, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., thymus, brain, and other tissue of the nervous system, blood cells, bone marrow, ovaries, and testes, and other reproductive tissue, mammary tissue, tonsils, melanocytes and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases of the thymus particularly thymic cancer and diseases involving T-cell maturation.

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# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 33

This gene is expressed primarily in human tonsils, and placenta, and to a lesser extent in adipocytes, melanocyte, and infant brain.

polypeptides are useful in providing immunological probes for differential identification gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an particularly of the inflammatory diseases, immune diseases, and obesity, expression of biological sample and for diagnosis of diseases and conditions: inflammatory diseases, immune diseases, and obesity. Similarly, polypeptides and antibodies directed to these of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, tissues and cell types (e.g., tonsils, placenta, adipocytes, melanocytes, and brain and fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard this gene at significantly higher or lower levels may be routinely detected in certain other tissue of the nervous system, and cancerous and wounded tissues) or bodily Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder. 2 22 30

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The tissue distribution and homology to this gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of diseases such as inflammation, immune diseases, and obesity.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 34

This gene is expressed in activated T cells, and to a lesser extent in pituitary, testis, and breast lymph node.

disorders of the immune system, expression of this gene at significantly higher or lower cells. Similarly, polypeptides and antibodies directed to these polypeptides are useful in levels may be routinely detected in certain tissues and cell types (e.g., pituilary, testes biological sample and for diagnosis of diseases and conditions: diseases relating to T and other reproductive tissue, mammary tissue, and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or providing immunological probes for differential identification of the tissue(s) or cell pinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in ype(s). For a number of disorders of the above tissues or cells, particularly, of the Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a nealthy tissue or bodily fluid from an individual not having the disorder. S 2 15

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of immune disorders.

# 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 35

This gene is expressed primarily in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the diseases relating to neurological disorders, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain, and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of neurological

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This gene is expressed primarily in infant brain

FEATURES OF PROTEIN ENCODED BY GENE NO: 36

S diseases relating to neurological disorders, expression of this gene at significantly providing immunological probes for differential identification of the tissue(s) or cel Similarly, polypeptides and antibodies directed to these polypeptides are useful in biological sample and for diagnosis of diseases and conditions: neurological disorders reagents for differential identification of the tissue(s) or cell type(s) present in a type(s). For a number of disorders of the above tissues or cells, particularly of the Therefore, polynucleotides and polypeptides of the invention are useful

2 5 standard gene expression level, i.e., the expression level in healthy tissue or bodily bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another brain and other tissue of the nervous system, and cancerous and wounded tissues) or higher or lower levels may be routinely detected in certain tissues and cell types (e.g., tissue or cell sample taken from an individual having such a disorder, relative to the

corresponding to this gene are useful for diagnosis and treatment of neurological The tissue distribution indicates that polynucleotides and polypeptides fluid from an individual not having the disorder.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 37

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This gene is expressed primarily in human ovary.

biological sample and for diagnosis of diseases and conditions: ovarian cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

type(s). For a number of disorders of the above tissues or cells, particularly of the providing immunological probes for differential identification of the tissue(s) or cel 25

expression of this gene at significantly higher or lower levels may be routinely detected in healthy tissue or bodily fluid from an individual not having the disorder. such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial in certain tissues and cell types (e.g., ovary and other reproductive tissue, and ovarian disorders such as those involving germ cells, ovarian follicles, stromal cells,

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corresponding to this gene are useful for diagnosis and treatment of ovariopathy The tissue distribution indicates that polynucleotides and polypeptides ઝ

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 38

reagents for differential identification of the tissue(s) or cell type(s) present in a This gene is expressed primarily in lymph node breast cancer. Therefore, polynucleotides and polypeptides of the invention are useful as

2 5 in healthy tissue or bodily fluid from an individual not having the disorder. expression of this gene at significantly higher or lower levels may be routinely detected polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: breast cancer. Similarly, such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial in certain tissues and cell types (e.g., mammary tissue and lymphoid tissue, and number of disorders of the above tissues or cells, particularly of the breast cancer, immunological probes for differential identification of the tissue(s) or cell type(s). For a

corresponding to this gene are useful for used as a diagnostic marker for breast cancer. The tissue distribution indicates that polynucleotides and polypeptides

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 39

25 20 30 biological sample and for diagnosis of diseases and conditions; neuronal disorders such differential identification of the tissue(s) or cell type(s). For a number of disorders of directed to these polypeptides are useful in providing immunological probes for as trauma, brain degeneration, and brain tumor. Similarly, polypeptides and antibodies reagents for differential identification of the tissue(s) or cell type(s) present in a another tissue or cell sample taken from an individual having such a disorder, relative to tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or significantly higher or lower levels may be routinely detected in certain tissues and cell the above tissues or cells, particularly of the brain, expression of this gene at the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. types (e.g., brain and other tissue of the nervous system, and cancerous and wounded Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in brain and to a lesser extent in other tissues.

ၾ corresponding to this gene are useful for diagnosis and therapeutic treatment of The tissue distribution indicates that polynucleotides and polypeptides

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 40

This gene is expressed in early stage human embryo, adrenal gland tumor, and immune tissues such as fetal liver, fetal spleen, T-cell, and myoloid progenitor cell line and to a lesser extent in ovary, colon cancer, and a few orther tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumorigenesis including adrenal gland tumor, colon cancer and various other tumors, developmental and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the cancer tissues, early stage human tissues, and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, spleen, blood cells, bone marrow, ovary

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in certain tissues and cell types (e.g., liver, spleen, blood cells, bone marrow, ovary and other reproductive tissue, and colon, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and therapeutic treatment of immune and developmental disorders, and tumorigenesis.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 41

This gene is expressed primarily in fetal lung, endothelial cells, liver, thymus and a few other immune tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as

oreagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune disorders such as immune deficiency and autoimmune diseases, pulmonary diseases, liver diseases, and tumor matasis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal lung, liver, endothelial cells, and immune tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain

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tissues and cell types (e.g., lung, endothelial cells, liver, thymus, and other tissue of the immune system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis of immune disorders and pulmonary and hepatic diseases. Its promoter may also be used for immune system and lung-

specific gene therapies. The expression of this gene in endothelial cells indicates that it

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nay also involve in angiogenesis which therefore may play role in tumor matasis.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 42

This gene is expressed primarily in liver, thyroid, parathyroid and to a lesser

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extent in fetal lung, stomach and early embryos.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: metabolic regulation, obesity, heptic failure, heptacellular tumors or thyroiditis and thyroid tumors. Similarly,

- polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive/endocrine system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, thyroid, parathyroid, lung, stomach, and embryonic tissue, and cancerous and wounded tissues) or bodily fluids
- stomach, and embryonic tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution and the extracellular locations indicates that polynucleotides and polypeptides corresponding to this gene are useful for the detection and treatment of digestive/endocrine disorders, including metabolic regulation, heptic failure, malabsortion, gastritis and neoplasms.

FEATURES OF PROTEIN ENCODED BY GENE NO: 43

cortex, hypothalmus and to a lesser extent in retina, adipose and stomach cancer and This gene is expressed primarily in Schizophrenic adult brain, pituitary, front

2 5 S of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells cell sample taken from an individual having such a disorder, relative to the standard or lower levels may be routinely detected in certain tissues and cell types (e.g., retinal particularly of the central nerve system, expression of this gene at significantly higher biological sample and for diagnosis of diseases and conditions: schizophrenia and other reagents for differential identification of the tissue(s) or cell type(s) present in a gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an tissue, adipose, stomach, and placenta, and cancerous and wounded tissues) or bodily polypeptides are useful in providing immunological probes for differential identification neurological disorders. Similarly, polypeptides and antibodies directed to these individual not having the disorder. fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or Therefore, polynucleotides and polypeptides of the invention are useful as

system, including schizophrenia, neurodegeneration, and neoplasia. Additionally, a corresponding to this gene are useful in treatment/detection of disorders in the nerve secreted protein in brain may serve as an endocrine. The tissue distribution indicates that polynucleotides and polypeptides

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 44

23 binding proteins which are thought to be important in signal transduction and protein transport. The translation product of this gene shares sequence homology with GTP

and to a lesser extent in gall bladder. cells, GM-CSF treated macrophage, anergic T cells, osteoblast, osteoclast, CD34+ cells This gene is expressed primarily in umbilical vein and microvascular endothelial

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35 immunological probes for differential identification of the tissue(s) or cell type(s). For a growth, osteonecrosis, osteoporosis, angiogenesis and/or hematopoeisis. Similarly, biological sample and for diagnosis of diseases and conditions: bone formation and reagents for differential identification of the tissue(s) or cell type(s) present in a hematopoeisis systems, expression of this gene at significantly higher or lower levels number of disorders of the above tissues or cells, particularly of the skeletal and polypeptides and antibodies directed to these polypeptides are useful in providing Therefore, polynucleotides and polypeptides of the invention are useful as

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expression level, i.e., the expression level in healthy tissue or bodily fluid from an sample taken from an individual having such a disorder, relative to the standard gene (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell cells, bone, and gall bladder, and cancerous and wounded tissues) or bodily fluids may be routinely detected in certain tissues and cell types (e.g., endothelial cells, blood individual not having the disorder.

hematopoeisis because its involvement in the growth signaling or angiogenesis. polynucleotides and polypeptides corresponding to this gene are useful for treatment/detection of bone formation and growth, osteonecrosis, osteoporosis, and/or The tissue distribution and homology to GTP binding proteins indicate that

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 45

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translocation on endoplasmic reticulum. sequence receptor gamma subunit which is thought to be important in protein The translation product of this gene shares sequence homology with signal

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to a lesser extent in endothelial cells and smooth muscle This gene is expressed primarily in adrenal gland, salivary gland, prostate, and

20 providing immunological probes for differential identification of the tissue(s) or cell biological sample and for diagnosis of diseases and conditions: protein secretion. secretory organs, expression of this gene at significantly higher or lower levels may be type(s). For a number of disorders of the above tissues or cells, particularly of the Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

25 ઝ routinely detected in certain tissues and cell types (e.g., adrenal gland, salivary gland, fluid from an individual not having the disorder prostate, endothelial cells, and smooth muscle, and cancerous and wounded tissues) or standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another

disorders, prostate cancer, xerostomia or sialorrhea polynucleotides and polypeptides corresponding to this gene are useful for endocrine The tissue distribution and homology to SSR gamma subunit indicate that

#### $\mathcal{Z}$ FEATURES OF PROTEIN ENCODED BY GENE NO: 46

melanocyte, amygdala, brain, and stomach. This gene is expressed primarily in osteoclastoma cells and to a lesser extent in

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: ossification, osteoporosis, fracture, osteonecrosis, osteosarcoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., melanocytes, amygdala, brain and other tissue of the nervous system,

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and stomach, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful in intervention of ossification, osteoporosis, fracture, osteonecrosis and osteosarcoma.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 48

20 The translation product of this gene shares sequence homology with proline rich proteins which is thought to be important in protein-protein interaction.

This gene is expressed primarily in brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurological and psychological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nerve system and endocrine system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded (issues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to proline-rich proteins indicate that polynucleotides and polypeptides corresponding to this gene are useful in intervention

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and detection of neurological diseases, including trauma, neoplasia, degenerative or metabolic conditions in the central nerve system. Additionally, the gene product may be a secreted by the brain as an endocrine.

# 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 49

The translation product of this gene shares sequence homology with the AOCB gene from Aspergillus nidulans which is important in asexual development.

This gene is expressed primarily in infant brain and to a lesser extent in the developing embryo, trachea tumors, B-cell lymphoma and synovial sarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurodegenerative diseases, leukemia and sarcoma's. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, blood cells, trachea, and synovial tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or

The tissue distribution in infant brain and sarcoma's and homology to a gene involved in a key step of eukaryotivc development (fungal spore formation) indicates that the protein product of this clone could play a role in neurological diseases such as schizophrenia, particularly in infants. The existence of the gene in a B-cell lymphoma indicates the gene may be used in the treatment and detection of leukernia.

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spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in

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healthy tissue or bodily fluid from an individual not having the disorder.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 50

30 This gene is expressed primarily in fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: pulmonary disorders including lung cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the pulmonary system, expression of this gene at significantly higher or

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in healthy tissue or bodily fluid from an individual not having the disorder. such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial lower levels may be routinely detected in certain tissues and cell types (e.g., lung, and

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pulmonary defects such as pulmonary edema and embolism, bronchitis and cystic the expression of this protein product in the adult could lead to lymphoma or sarcoma role in development of the pulmonary system. This would suggest that misregulation of formation, particularly in the lung. It may also be involved in predisposition to certain The tissue distribution of this gene only in fetal lung indicates that it plays a key

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 51

This gene is expressed primarily in hematopoietic cell types and fetal cells and to

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a lesser extent in all tissue types reagents for differential identification of the tissue(s) or cell type(s) present in a system and hematopoeisis. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: defects in the immune Therefore, polynucleotides and polypeptides of the invention are useful as

of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another particularly of the immune and hematopoietic systems, expression of this gene at standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the types (e.g., hematopoietic cells, and fetal tissue, and cancerous and wounded tissues) significantly higher or lower levels may be routinely detected in certain tissues and cel

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8 the developing embryo indicates that polynucleotides and polypeptides corresponding to affecting the immune system or hematopoeisis disorders such as leukemia, AIDS this gene are useful for detection and treatment of lymphomas and disease states arthritis and asthma. The tissue distribution of this gene predominantly in hematopoeitic cells and in

fluid from an individual not having the disorder.

#### ႘ FEATURES OF PROTEIN ENCODED BY GENE NO: 52

fetal liver, infant brain and T cell leukemias. This gene is expressed primarily in prostate and to a lesser extent in fetal spleen

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significantly higher or lower levels may be routinely detected in certain tissues and cell particularly of the immune system, and/or prostate gland expression of this gene at of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification prostate cancer, leukemia. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions; prostate disorders reagents for differential identification of the tissue(s) or cell type(s) present in a types (e.g., thymus, spleen, liver, brain and other tissue of the nervous system, and Therefore, polynucleotides and polypeptides of the invention are useful as

blood cells, and cancerous and wounded tissues) or bodily sluids (e.g., serum, plasma urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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polypeptides corresponding to this gene are useful for detection or treatment of prostate may play a role in the immune system and its misregulation could lead to immune disorders or prostate cancer. Its distribution in fetal liver and fetal spleen indicates it disorders such as leukemia, arthritis and asthma. The tissue distribution of this gene in prostate indicates that polynucleotides and

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 53

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This gene is expressed primarily in brain. The translation product of this gene shares sequence homology with dynein.

reagents for differential identification of the tissue(s) or cell type(s) present in a of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identificatior diseases of the brain. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: neuro-degenerative Therefore, polynucleotides and polypeptides of the invention are useful as

- છ gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an and other tissue of the nervous system, and cancerous and wounded tissues) or bodily or lower levels may be routinely detected in certain tissues and cell types (e.g., brain particularly neuro-degenerative diseases expression of this gene at significantly higher cell sample taken from an individual having such a disorder, relative to the standard fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or
- 35 individual not having the disorder

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molecules indicates that polynucleotides and polypeptides corresponding to this gene are useful for detection/treatment of neurodegenerative diseases, such as Alzheimers, The predominant tissue distribution in the brain and homology to dynein, a microtubule motor protein involved in the positioning of cellular organelles and Huntigtons, Parkinsons diseases and shizophrenia.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 54

The translation product of this gene shares sequence homology with ubiquitinconjugation protein, an enzyme which is thought to be important in the processing of the Huntingtons Disease causing gene.

This gene is expressed primarily in brain and to a lesser extent in activated

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

such a disorder, relative to the standard gene expression level, i.e., the expression level cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial significantly higher or lower levels may be routinely detected in certain tissues and cell disease states including Huntington's disease. Similarly, polypeptides and antibodies fluid or spinal fluid) or another tissue or cell sample taken from an individual having biological sample and for diagnosis of diseases and conditions: neurodegenerative differential identification of brain tissues. For a number of disorders of the above tissues or cells, particularly of the neurological systems expression of this gene at directed to these polypeptides are useful in providing immunological probes for types (e.g., brain and other tissue of the nervous system, and blood cells, and 2 8

The predominant tissue distribution of this gene in the brain and its homology to a Huntington interacting protein indicates that polynucleotides and polypeptides corresponding to this gene are useful for the regulation of the expression of the in healthy tissue or bodily fluid from an individual not having the disorder. Huntington disease gene and other neurodegenerative diseases including

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muscular atrophy. In addition, the existence of elevated levels of free ubiquitin pools in Alzheimer's disease, Parkinson's disease and amylotrophic lateral sclerosis indicates Thus, considering the gene described here is homologous to a ubiquitin-conjugation that the ubiquitin pathway of protein degradation plays a role in these disease states. spinocerebullar ataxia types I and III, dentatorubropallidoluysian and spinal bulbar protein it may play a general role in neurodegenarative conditions. 33

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# FEATURES OF PROTEIN ENCODED BY GENE NO: 56

This gene is expressed primarily in T-cells (anergic T-cells, resting|T-Cells, apoptotic T-cells) and lymph node (breast), as well as brain (hypothalamus hippocampus, pituitary, infant brain, early-stage brain).

immunological probes for differential identification of the ussue(s) or cell type(s). For a cells, lymphoid tissue, and brain and other tissue of the nervous system, and cancerous neurological (e.g. Alzheimer's disease, dementia, schizophrenia) disorders. Similarly, hematopoietic and immune systems, expression of this gene at significantly higher or number of disorders of the above tissues or cells, particularly of the central nervous, and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in immunodeficiencies, autoimmunities, inflammation, leukemias & lymphomas) and lower levels may be routinely detected in certain tissues and cell types (e.g., blood Therefore, polynucleotides and polypeptides of the invention are useful as polypeptides and antibodies directed to these polypeptides are useful in providing reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune (e.g. healthy tissue or bodily fluid from an individual not having the disorder. 2 2 S

associated with the hematopoietic and immune systems, such as anemias (leukemias). developmental brain defects, neuro-degenerative diseases or behavioral abnomalities corresponding to this gene are useful in the intervention or detection of pathologies The tissue distribution indicates that polynucleotides and polypeptides In addition, the expression in brain (including fetal) might suggest a role in (e.g. schizophrenia, Alzheimer's, dementia, depression, etc.). ಜ

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 57

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This gene is expressed primarily in lung, and to a lesser extent in a variety of other hematological cell types (e.g. Raji cells, bone marrow cell line, activated monocytes).

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polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vasculo-pulmonary and hematopoietic systems, expression of this hematological disfunction. Similarly, polypeptides and antibodies directed to these Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: pulmonary and/or reagents for differential identification of the tissue(s) or cell type(s) present in a

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standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another and cell types (e.g., lung and blood cells, and cancerous and wounded tissues) or gene at significantly higher or lower-levels may be routinely detected in certain tissues tissue or cell sample taken from an individual having such a disorder, relative to the

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in a variety of leukocytic cell types and a bone marrow cell line might suggest a role in inflammation, immunodeficiencies and autoimmunities. hematopoietic and immune system disorders, such as leukemias & lymphomas, associated with the vasculo-pulmonary system. In addition the expression of this gene corresponding to this gene are useful in the intervention and detection of pathologies The tissue distribution indicates that polynucleotides and polypeptides

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 58

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the standard gene expression level, i.e., the expression level in healthy tissue or bodily

2 ADP in the presence of ATP or inorganic triphosphate. taurus]), which is thought to be important in catalyzing the phosphorylation of AMP to kinase isozyme 3 (gil163528 GTP:AMP phosphotransferase (EC 2.7.4.10) [Bos The translation product of this gene shares sequence homology with adenylate

extent in many other tissues This gene is expressed primarily in fetal liver, heart and placenta, and to a lesser

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hematapoiesis, as well as immunological disorders.

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of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification or reproductive disorders. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: hepatic, cardiovascular reagents for differential identification of the tissue(s) or cell type(s) present in a bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another and cell types (e.g., liver, heart, and placenta, and cancerous and wounded tissues) or gene at significantly higher or lower levels may be routinely detected in certain tissues particularly of the hepatic, cardiovascular and reproductive systems, expression of this tissue or cell sample taken from an individual having such a disorder, relative to the Therefore, polynucleotides and polypeptides of the invention are useful as

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corresponding to this gene are useful for the treatment and diagnosis of conditions development and the differentiation of hepatocyte progenitor cells. related to hepatic function and pathogenesis, in particular, those dealing with liver The tissue distribution indicates that polynucleotides and polypeptides

standard gene expression level, i.e., the expression level in healthy tissue or bodily

fluid from an individual not having the disorder.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 59

Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in CD34 positive cells (Cord Blood).

- 5 reagents for differential identification of the tissue(s) or cell type(s) present in a these polypeptides are useful in providing immunological probes for differential differentiation and immune disorders. Similarly, polypeptides and antibodies directed to biological sample and for diagnosis of diseases and conditions: hematopoietic another tissue or cell sample taken from an individual having such a disorder, relative to tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or and cell types (e.g., hematopoietic cells, and blood cells, and cancerous and wounded gene at significantly higher or lower levels may be routinely detected in certain tissues tissues or cells, particularly of hematopoietic and immune systems, expression of this identification of the tissue(s) or cell type(s). For a number of disorders of the above
- associated with CD34-positive cells, and therefore as a marker for cell differentiation in corresponding to this gene are useful in the detection and treatment of conditions fluid from an individual not having the disorder. The tissue distribution indicates that polynucleotides and polypeptides

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 60

sequence identity to the murine gene designated Insulin-Like Growth Factor-Binding The translation product of the predicted open reading frame of this contig has

25 (1994)Protein (IGFBP)-1 as described by Lee and colleagues (Hepatology 19 (3), 656-665

This gene is expressed exclusively in hemangiopericytoma.

reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

- biological sample and for diagnosis of hemangiopericytoma and other pericyte or endothelial cell proliferative disorders. Similarly, polypeptides and antibodies directed gene at significantly higher or lower levels may routinely be detected in certain tissues tissues or cells, particularly of the circulatory and immune systems, expression of this identification of the tissue(s) or cell type(s). For a number of disorders of the above to these polypeptides are useful in providing immunological probes for differential
- 35 and cell types (e.g., pericyte or endothelial cells, and liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

Polynucleotides and polypeptides corresponding to this gene are useful as cell growth regulators since IGFBP-1-like molecules function as modulators of insulin-like growth factor activity. In addition, since IGFBP-1 is expressed at high levels following hepatectomy and during fetal liver development, polynucleotides of the present invention may also be used for the diagnosis of developmental disorders. Further, polypeptides of the present invention may be used therapeutically to treat developmental liver disorders as well as to regulate hepatocyte and supporting cell growth following hepatectomy or to treat liver disorders.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hemangiopericytoma and liver disorders.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 61

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This gene is expressed primarily in schizophrenic frontal cortex.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: nervous system and cognitive disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the frontal cortex and CNS expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, treatment and diagnosis of frontal cortex, neuro-degenerative and CNS disorders

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# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 62

This gene is expressed primarily in human adrenal gland tumor, and to a lesser extent in human kidney, medulla and adult pulmonary tissue.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: metabolic endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are

- suseful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine and nervous system disorders and neoplasia, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., adrenal gland, kidney, brain and other tissue of the nervous system, pulmonary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum,
- 0 pulmonary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- 15 The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, treatment and diagnosis of neurological and endocrine disorders including neoplasia.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 63

This gene is expressed primarily in human adipocytes, and to a lesser extent in spleen, 12-week old human, and testes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune, metabolic and growth disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types(e.g., adipocytes,

- 30 spleen, and testes and other reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- 35 The tissue distribution indicates that polynucleotides and polypeplides corresponding to this gene are useful for study, diagnosis and treatment of immune, developmental and metabolic disorders.

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FEATURES OF PROTEIN ENCODED BY GENE NO: 64
One translated product of this clone is homologous to the mouse zinc finger protein
PZF. (See Accession No. 453376; see also Gene 152 (2), 233-238 (1995).) Preferred

- 5 polypeptide fragments correspond to the highly conserved domains shared between mouse and man. For example, preferred polypeptide fragments comprise the amino acid sequence: LQCEICGFTCRQKASLNWHMKKHDADSFYQFSCNICGKKFEKKDSVVAHKAKSH PEV (SEQ ID NO: 621); ITSTDILGTNPESLTQPSD (SEQ ID NO: 622); NSTSGECLLLEAEGM SKSY (SEQ ID NO: 623); CSGTERVSLMADGKIFVGSGSSGGTEGLVMNSDILGATTEVLIEDSD SAGP (SEQ ID NO: 624); IQYVRCEMEGCGTVLAHPRYLQHHIKYQHLLKKKYVCPHPSCGRLF RLQKQLLRHAKHHT (SEQ ID NO: 625); DQRDYICEYCARAFKSSHNLAVHRMIHTGEK (SEQ ID NO: 626); RSSRTSVSRHRDTENTRSSRSKTGSLQLICKSEPNTDQLDY (SEQ ID NO: 627); PFKDDPRDETYKPHLERETPKPRRKSG (SEQ ID NO: 630); QYVRCEMEGCGTVLAHPRYLQ HHIKYQHLLKKKYVCPHPSCGRLFRLQKQLLRHAKHHTD (SEQ ID NO: 629); or residues
- 15 151-182 of QRDYICEYCARAFKSSHNLAVHRMIHTGEKHY (SEQ ID NO: 628). Also preferred are polynucleotide fragments encoding these polypeptide fragments.

This gene is expressed primarily in Rhabdomyosarcoma, melanocyte and colon cancer tissue and to a lesser extent in smooth muscle, pancreatic tumor, and apoptotic T-cells.

- Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to.. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of
- or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and hemopoetic, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., striated muscle, melanocytes, colon, smooth muscle, pancreas, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, diagnosis and treatment of cancer and becomes discorder.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 65

This gene is expressed primarily in human adipose and salivary gland tissue and to a lesser extent in human bone marrow and fetal kidney.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

- biological sample and for diagnosis of diseases and conditions: metabolic and immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are disorders. Similarly, polypeptides and antibodies directed to these polypeptides are disorders. Similarly, polypeptides and antibodies directed to these polypeptides are disorders. Similarly, polypeptides and antibodies directed to these polypeptides are disorders of the above tissues or cells, particularly of or cell type(s). For a number of disorders of the above tissues or cells, particularly of the metabolic and hemopoetic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., adipose, lower levels may be routinely detected in certain tissues and wounded tissues) or salivary gland, bone marrow, and kidney, and cancerous and wounded tissues) or salivary gland, bone marrow, and kidney, and cancerous and wounded tissues) or solily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily
- 15 standard gene expression of the disorder.

  fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, diagnosis of metabolic and immune disorders.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 66

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This translated product of this gene was recently identified as oxytocinase splice variant This translated product of this gene was recently identified as oxytocinase splice variant 1. (See Accession Nos. 2209276 and d1010078.) Preferred polypeptide fragments 1. (See Accession Nos. 2209276 and d1010078.) Preferred polypeptide fragments comprise the amino acid sequence: EMFDSLSYFKGSSLLLMLKTYLSEDVFQHAVVLYLHN comprise the amino acid sequence: EMFDSLSYFKGSSLLLMLKTYLSEDVFQHAVVLYLHN COMPRISE THE ACCOUNTY OF THE

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 67

This gene is expressed primarily in hemopoetic cells, particularly apoptotic T-cells, and to lesser extent in primary dendritic cells and adipose tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of apoptotic T-cells, primary dentritic cells, and adipose tissue present in a biological sample and for diagnosis of diseases and adipose tissue present in a biological sample and general immune disorders.

reagents for differential identification of apopular and for diagnosis of diseases and adipose tissue present in a biological sample and for diagnosis of diseases and conditions; hemopoetic diseases including cancer and general immune disorders.

Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell

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type(s). For a number of disorders of the above tissues or cells, particularly of the oral and intestinal mucosa as well as hemopoetic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of diseases of the immune system, including cancer, hemopoetic and infectious diseases.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 68

This gene is expressed primarily in kidney cortex and to a lesser extent in infant

15 brain, heart, uterus, and blood.
Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of kidney tissue present in a biological sample and for diagnosis of diseases and conditions: soft tissue cancer, inflammation, kidney fibrosis. Similarly, polypeptides and antibodies directed to these polypeptides are useful

- type(s). For a number of disorders for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and endocrines systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., kidney, brain, and other nervous tissue, heart, uterus, and blood cells, and cancerous and
- orain, and outer nervous ussue, neads, are us, and other cents, and cancerous and 25 wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides.

30 corresponding to this gene are useful for study and treatment of cancer and fibroses.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 69

The translation product of this gene shares strong sequence homology with vertebrate and invertebrate protein tyrosine phosphatases.

This gene is expressed primarily in endometrial tumors, melanocytes, myeloid progenitors and to a lesser extent in infant brain, adipocytes, and several hematopoietic stem cells.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of transformed hematopoietic and epithelial cells present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, of skin and endometrium, leukemia. Similarly,

- polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and hemopoietic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium; melanocytes, bone marrow, adipocytes, hematopoietic cells, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the
- The tissue distribution and sequence similarity with tyrosine phosphatases indicate that polynucleotides and polypeptides corresponding to this generare useful for study and treatment of cancer and hematopoietic disorders.

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# 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 70

This gene is expressed primarily in osteoclastoma, breast, and infant brain and to a lesser extent in various fetal and transformed bone, ovarian, and neuronal cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

- biological sample and for diagnosis of diseases and conditions: degenerative conditions of the brain and skeleton. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and skeletal system, expression of this gene at significantly on higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone, mammary tissue, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having
- 35 in healthy tissue or bodily fluid from an individual not having the disorder.

such a disorder, relative to the standard gene expression level, i.e., the expression level

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study and treatment of degenerative, neurological and skeletal disorders.

# 5 FEATURES OF PROTEIN: ENCODED BY GENE NO: 71

This gene was originally cloned from tumor cell lines. Recently another group has also cloned this gene, calling it the human malignant melanoma metastasis-suppressor (KiSS-1) gene. (See Accession No. U43527.) Preferred polypeptide fragments comprise the amino acid sequence: LEKVASVGNSRPTGQQLESLGLLA (SEQ ID NO: 632); VHREEASCYCQAEPSGDL (SEQ ID NO: 633); RPALRQAGGGTREPRQKRWAGL (SEQ ID NO: 634); and AVNFRPQRSQSM (SEQ ID NO: 635). Any frame shifts can easily be resolved using known molecular biology techniques.

This gene is expressed primarily in many types of carcinomas and to a lesser extent in many normal organs.

- Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissues(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer particularly melanomas, and other hyperproliferative disorders.

  Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell
- providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of transformed organ tissue, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another rissue or cell sample taken from an individual having such a disorder, relative to
- another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. As a tumor suppressor gene, increase amounts of the polypeptide can be used to treat patients having a particular cancer.

The tissue distribution indicates that this gene and the translated product is useful for diagnosing and study of cancer.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 72

This gene is expressed primarily in striatum and to a lesser extent in adipocytes and hemangioperiocytoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of striatal cells present in a biological sample and for diagnosis of diseases and conditions: neurological, fat and lysosomal storage

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diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., striatal tissue, adipocytes, and vascular tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis, study and treatment of neurodegenerative and growth disorders.

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individual not having the disorder.

# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 73

This gene is expressed primarily in bone marrow stromal cells and to a lesser extent in smooth muscle, testes, endothelium, and brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of bone marrow present in a biological sample and 20 for diagnosis of diseases and conditions: connective tissue and hematopoietic diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal and hematopoietic systems, expression of this gene at significantly higher or

25 lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, stromal cells, smooth muscle, testes and other reproductive tissue, endothelium, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to 30 the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, diagnosis, and treatment of connective tissue and blood diseases.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 74

This gene is expressed primarily in brain, fetal liver and lung and to a lesser extent in retina, spinal chord, activated T-cells and endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of brain and regenerating liver present in a biological sample and for diagnosis of diseases and conditions: CNS and spinal chord injuries, immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, liver, pulmonary tissue, blood cells, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study and treatment of hematopoietic and neurological conditions.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 75

The translation product of this gene shares sequence homology with GTP binding proteins (intracellular).

This gene is expressed primarily in bone marrow, brain, and melanocytes and to a lesser extent in various endocrine and hematopoietic tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: hematopietic and nervous system conditions. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous and immune, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, melanocytes, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder,

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relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to nucleotide binding factors indicate that polynucleotides and polypeptides corresponding to this gene are useful for study,

5 diagnosis, and treatment of brain degenerative, skin and blood diseases.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 76

This gene is expressed primarily in activated T-cells and to a lesser extent in retina, brain, and fetal bone.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of activated T-cells and developing brain present in a biological sample and for diagnosis of diseases and conditions: immune deficiencies and skeletal and neuronal growth disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous, immune, and skeletomuscular sustems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, brain and other tissue of the nervous system, retinal tissue, and bone, and cancerous and wounded tissues) or

bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis, study and treatment of cancer, urogenital, and brain degenerative diseases.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 77

This gene is expressed primarily in fetal liver, activated monocytes, osteoblasts and to a lesser extent in synovial, brain, and lymphoid tissues.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of myeloid and lymphoid present in a biological sample and for diagnosis of diseases and conditions: inflammation, immune deficiencies, cancer. Similarly, polypeptides and antibodies directed to these

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and skeleton, expression of this gene at significantly

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higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, blood cells, bone, synovial tissue, brain and other tissue of the nervous system, and lymphoid tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for study, diagnosis, and treatment of lymphoid and marginal diseases.

10 and mesenchymal cancers and nervous system diseases.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 7

The translation product of this gene shares sequence homology with polymerase polyprotein precursor which is thought to be important in DNA repair and replication or the case is consecred primarily in infant brain and to a lesser extent in tumors.

This gene is expressed primarily in infant brain and to a lesser extent in tumors and tumor cell lines

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, especially of the neural system and developing organs. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neural system expression of this gene at significantly higher or lower levels may be routinely detected

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in certain (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to polymerase polyprotein precursor indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancers especially of the neural system and developing organs

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# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 79

This gene is expressed primarily in muscle and endothelial cells and to a lesser extent in brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: vascular diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain (e.g., muscle, endothelial cells, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum,

10 plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e. the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution indicates that polynucleotides and polypeptides
corresponding to this gene are useful for treatment and diagnosis of disorders of the
vascular and neural system including cardiovascular and endothelial.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 80

This gene is expressed primarily in placenta and to a lesser extent in fetal liver
Therefore, polynucleotides and polypeptides of the invention are useful as

25 20 30 immunological probes for differential identification of the tissue(s) or cell type(s). For a reagents for differential identification of the tissue(s) or cell type(s) present in a of this gene at significantly higher or lower levels may be routinely detected in certain number of disorders of the above tissues or cells, particularly of developmental and disorder of the haemopoietic system, fetal liver and placenta. Similarly biological sample and for diagnosis of diseases and conditions: developmental disorders bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another polypeptides and antibodies directed to these polypeptides are useful in providing standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the tissues and cell types (e.g., placenta and liver, and cancerous and wounded tissues) or disorders and disorder of the haemopoietic system, fetal liver and placenta, expression Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of developmental disorders and disorders of the haemopoietic system, fetal liver and placenta.

fluid from an individual not having the disorder.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 81

This gene is expressed primarily in bone marrow, placenta and tissues and organs of the hematopoietic system.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: disorders of the bone and haemopoietic system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, bone and hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone marrow, placenta, and hematopoietic cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorders of the income.

20 immune, bone and hematopoietic system

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 82

The translation product of this gene shares sequence homology with secretory carrier membrane protein which is thought to be important in protein transport and export. Any frame shifts in coding sequence can be easily resolved using standard molecular biology techniques. Another group recently cloned this gene, calling it SCAMP. (See Accession No. 2232243.)

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This gene is expressed primarily in prostate, breast and spleen, and to a lesser extent in several other tissues and organs.

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: disorders of the breast prostate and spleen. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly disorders of the breast prostate and spleen, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell

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types (e.g., prostate, mammary tissue, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to secretory carrier membrane protein indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of disorders of the breast, prostate and spleen.

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# 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 83

This gene is expressed primarily in developing organs and tissue like placenta and infant brain and to a lesser extent in developed organs and tissue like cerebellum and heart.

routinely detected in certain tissues and cell types (e.g., placenta, heart, brain and other biological sample and for diagnosis of diseases and conditions: neurological diseases. sample taken from an individual having such a disorder, relative to the standard gene neural system, expression of this gene at significantly higher or lower levels may be providing immunological probes for differential identification of the tissue(s) or cell issue of the nervous system, and cancerous and wounded tissues) or bodily fluids expression level, i.e., the expression level in healthy tissue or bodily fluid from an type(s). For a number of disorders of the above tissues or cells, particularly of the (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell Similarly, polypeptides and antibodies directed to these polypeptides are useful in Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder. 15 ន 25

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of diseases of the neural system including neurological disorders and cancer.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 84

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The translation product of this gene shares sequence homology with ATPase 6 in Trypanosoma brucei which is thought to be important in metabolism.

This gene is expressed primarily in tumor and fetal tissues and to a lesser extent in melanocytes, kidney cortex, monocytes and ovary.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions: metabolism disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., fetal tissues, melanocytes, kidney, blood cells, ovary and other tissue of the reproductive system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

The tissue distribution and homology to ATPase indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of metabolism disorders, especially in fetal and tumor tissue growth.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 85

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fluid from an individual not having the disorder

The translation product of this gene shares sequence homology with the immunoglobulin superfamily of proteins which are known to be important in immune response and immunity.

20 This gene is expressed primarily in stromal cells, colon cancer, lung, amygdala, melanocyte and to a lesser extent in a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of stromal cell development and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the stromal cells, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., stromal cells,

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colon, lung, amygdala, and melanocytes, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to immunoglobulin indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of immune system disorders.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 86

The translation product of this gene shares sequence homology with transcription iniation factor eIF-4 gamma which is thought to be important in gene transcription

This gene is expressed primarily in tumor tissues

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumorigenesis.

- providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly in tumor tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution and homology to transcription iniation factor eIF-4
  gamma indicate that polynucleotides and polypeptides corresponding to this gene are
  useful for gene regulation in tumorigenesis.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 87

The translation product of this gene shares sequence homology at low level in prolines with secreted basic proline-rich peptide II-2 which is thought to be important in protein structure or inhibiting hydroxyapatite formation in vitro.

This gene is expressed primarily in endometrial tumor and fetal lung.

Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of the tissue(s) or cell type(s) present in a

biological sample and for diagnosis of diseases and conditions: endometrial tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the muscular/skeletal and reproductive systems, expression of this gene at significantly

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35 higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endometrium, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample

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taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to secreted basic proline-rich peptide II-2 indicate that polynucleotides and polypeptides corresponding to this gene are useful for inhibiting hydroxyapatite formation or establishing cell/tissue structure.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 88

This gene is expressed primarily in: amniotic cells inducted with TNF in culture; and to a lesser extent in colon tissue from a patient with Crohn's Disease; parathyroid tumor; activated T-cells; cells of the human Caco-2 cell line; adenocarcinoma; colon; corpus colosum; fetal kidney; pancreas tumor; fetal brain; early stage brain, and anergic

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system;

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c.g., tumors, expression of this gene at significantly higher or lower levels may be routinely detected in certain (e.g., amniotic cells, colon, kidney, pancreas, parathyroid, brain and other tissue of the nervous system, blood cells, hematopoietic cells, liver, spleen, bone, testes and other reproductive tissue, brain and other tissue of the nervous system, and epithelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product of this clone is useful for modulating tumorigenesis and other immune system conditions such as disorders in immune response.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 89

This gene is expressed primarily in fetal liver/spleen and hematopoietic cells and

to a lesser extent in brain, osteosarcoma, and testis tumor.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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biological sample and for diagnosis of diseases and conditions: leukemia and hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

significantly of the hematopoietic and immune systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., hematopoietic cells, liver, spleen, bone, testes, and other reproductive tissue, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of hematopoietic and immune disorders.

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fluid from an individual not having the disorder.

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 90

The translation product of this gene shares weak sequence homology with mouse Gcap1 protein which is developmentally regulated in brain.

This gene is expressed primarily in infant and adult brain and fetal liver/spleen and to a lesser extent in smooth muscle, T cells, and a variety of other tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neurological or hematopoietic disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous, hematopoietic, immune, and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain

itssues and cell types (e.g., brain and other tissue of the nervous system, blood cells, liver, spleen, and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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diagnosis of disorders in neuronal, hematopoietic, immune, and endocrine systems. polynucleotides and polypeptides corresponding to this gene are useful for treatubg and The tissue distribution and its homology to Gcap1 protein indicate that

#### S FEATURES OF PROTEIN ENCODED BY GENE NO: 91

extent in tumor tissues This gene is expressed primarily in brain and hematopoietic cells and to a lesser

reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

- 5 biological sample and for diagnosis of diseases and conditions; disorder in nervous, of the above tissues or cells, particularly of the in nervous, hematopoietic, immune antibodies directed to these polypeptides are useful in providing immunological probes hematopoietic, immune systems and tumorigenesis. Similarly, polypeptides and for differential identification of the tissue(s) or cell type(s). For a number of disorders
- 20 5 system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, systems, expression of this gene at significantly higher or lower levels may be routinely urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an detected in certain tissues and cell types (e.g., brain and other tissue of the nervous the expression level in healthy tissue or bodily fluid from an individual not having the individual having such a disorder, relative to the standard gene expression level, i.e.,
- systems. for diagnosis and treatment of disorders in the nervous, hematopoietic, and immune The tissue distribution indicates that the protein product of this clone is useful

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 92

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fluid or spinal fluid) or another tissue or cell sample taken from an individual having

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neuroendocrine-specific protein A which is thought to be important in neurologic The translation product of this gene shares sequence homology with

30 This gene is expressed primarily in brain tissues

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neural disorders and degeneration disease. Similarly, polypeptides and antibodies directed to these Therefore, polynucleotides and polypeptides of the invention are useful as

35 of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification particularly of the central or peripheral nervous systems, expression of this gene at

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such a disorder, relative to the standard gene expression level, i.e., the expression level types (e.g., hematopoietic cells, and brain and other tissue of the nervous system, and fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial significantly higher or lower levels may be routinely detected in certain tissues and cell

in healthy tissue or bodily fluid from an individual not having the disorder The tissue distribution and homology to neuroendocrine-specific protein A

treatment or diagnosis of neural disorders and degeneration disease. indicate that polynucleotides and polypeptides corresponding to this gene are useful for

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 93

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like protein and prolin-rich protein which are thought to be important in connective tissue function and tissue structure. The translation product of this gene shares sequence homology with collagen-

20 5 polypeptides are useful in providing immunological probes for differential identification cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, biological sample and for diagnosis of diseases and conditions: neuronal or reagents for differential identification of the tissue(s) or cell type(s) present in a significantly higher or lower levels may be routinely detected in certain tissues and cell particularly of the nervous and hematopoietic systems, expression of this gene at hematopoietic disorders. Similarly, polypeptides and antibodies directed to these types (e.g., liver, spleen, and brain and other tissue of the nervous system, and This gene is expressed primarily in fetal liver/spleen and brain tissues. Therefore, polynucleotides and polypeptides of the invention are useful as

30 useful for supporting brain and hematopoietic tissue function and diagnosis and proteins indicate that polynucleotides and polypeptides corresponding to this gene are in healthy tissue or bodily fluid from an individual not having the disorder. such a disorder, relative to the standard gene expression level, i.e., the expression level The tissue distribution and homology to collagen-like protein and proline-rich

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 94

treatment of disorders in these functions

35 reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as This gene is expressed primarily in embryonic tissues and tumor tissues.

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biological sample and for diagnosis of diseases and conditions which include, but are not limited to. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system (e.g., tumors), expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancer.

# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 95

This gene is expressed primarily in brain tumor, placenta, and melanoma.

Therefore, polynucleotides and polypeptides of the invention are useful as

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: brain tumor or melanoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain or melanocytes, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, placenta, and melanocytes, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

30 The tissue distribution indicates that the translation product of this gene is useful in the diagnosis and treatment of brain tumors and melanoma.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 96

The translation product of this gene shares sequence homology with a yeast membrane protein, SUR4, which encodes for APA1 that acts on a glucose-signaling pathway that controls the expression of several genes that are transcriptionally regulated by glucose.

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This gene is expressed primarily in fetal liver, and to a lesser extent in placenta

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of fetal liver or defects of glucose-regulated ATPase activities in tissues. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal immune/hematopoietic system,

in certain tissues and cell types (e.g., liver, placenta, and mammary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual or having the disorder.

The tissue distribution and homology to yeast SUR4 membrane protein indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of defects of fetal liver or defects of glucose-regulated ATPase

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 97

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This gene is expressed primarily in fetal liver, brain, and amniotic fluid.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of the fetal immune system and adult brain. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal immune system and adult brain, expression of this gene at

significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., amniotic fluid, serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder

The tissue distribution indicates that the protein product of this clone is useful for detecting defects of the fetal immune and hematopoietic systems since fetal liver is

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product of this gene is thought to be useful for detecting certain neurological defects of the predominant organ responsible for hematopoiesis in the fetus. In addition, the gene

#### S FEATURES OF PROTEIN ENCODED BY GENE NO: 98

protein precursor, Vitellogenin which is thought to be important in binding lipids such The translation product of this gene shares sequence homology with an yolk

This gene is expressed primarily in amniotic cells and fetal liver

5 20 5 biological sample and for diagnosis of diseases and conditions: defects in amniotic for differential identification of the tissue(s) or cell type(s). For a number of disorders cells, fetal liver development and the fetal immune system. Similarly, polypeptides and reagents for differential identification of the tissue(s) or cell type(s) present in a urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an and liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, levels may be routinely detected in certain tissues and cell types (e.g., amniotic cells, state is likely, e.g., immunel, expression of this gene at significantly higher or lower of the above tissues or cells, particularly of the [insert system where a related disease antibodies directed to these polypeptides are useful in providing immunological probes the expression level in healthy tissue or bodily fluid from an individual not having the individual having such a disorder, relative to the standard gene expression level, i.e., Therefore, polynucleotides and polypeptides of the invention are useful as

25 product of this clone is useful for treatment and diagnosis of defects in amniotic cells fetal liver development and the fetal immune system. The tissue distribution and homology to vitellogenin indicate that the protein

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 99

and stromal cells This gene is expressed primarily in placenta, endometrial tumor, osteosarcoma

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biological sample and for diagnosis of diseases and conditions: tumor of the reagents for differential identification of the tissue(s) or cell type(s) present in a the above tissues or cells, particularly of the obstetric system (e.g. placenta, differential identification of the tissue(s) or cell type(s). For a number of disorders of directed to these polypeptides are useful in providing immunological probes for endometrium or bone, and osteosarcoma. Similarly, polypeptides and antibodies Therefore, polynucleotides and polypeptides of the invention are useful as

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cell sample taken from an individual having such a disorder, relative to the standard endometrium, bone, and stromal cells, and cancerous and wounded tissues) or bodily individual not having the disorder gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or levels may be routinely detected in certain tissues and cell types (e.g., placenta, endometrium) and the bones, expression of this gene at significantly higher or lower

5 aforementioned tissues.. abnormalities of the endometrium, and the bones because of its abundance in the corresponding to this gene are useful for diagnosis and treatment of turnors and The tissue distribution indicates that polynucleotides and polypeptides

#### FEATURES OF PROTEIN ENCODED BY GENE NO:

This gene is expressed primarily in hepatocellular tumor

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23 type(s). For a number of disorders of the above tissues or cells, particularly of the liver biological sample and for diagnosis of diseases and conditions: hepatocellular tumor standard gene expression level, i.e., the expression level in healthy tissue or bodily in certain tissues and cell types (e.g., liver, and cancerous and wounded tissues) or providing immunological probes for differential identification of the tissue(s) or cell Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another expression of this gene at significantly higher or lower levels may be routinely detected fluid from an individual not having the disorder tissue or cell sample taken from an individual having such a disorder, relative to the Therefore, polynucleotides and polypeptides of the invention are useful as

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for diagnosis and treatment of hepatocellular cancer because of its abundant expression in this tissue The tissue distribution indicates that the protein product of this clone is useful

## FEATURES OF PROTEIN ENCODED BY GENE NO: 101

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This gene is expressed primarily in Corpus Colosum, fetal lung and infant

brain

35 reagents for differential identification of the tissue(s) or cell type(s) present in a Colosum or defects of the fetal lung. Similarly, polypeptides and antibodies directed to biological sample and for diagnosis of diseases and conditions: defects of the Corpus Therefore, polynucleotides and polypeptides of the invention are useful as

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these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the Corpus Colosum and brain in general, and fetal lung, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., lung, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of defects of the Corpus Colosum and brain in general, and defects of fetal lung.

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# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 102

This gene is expressed primarily in T cells and stromal cells, and to a lesser extent in adrenal gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of T cell immunity and stromal cell development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, stromal cells, and adrenal gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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fluid from an individual not having the disorder.

The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of defects of T cell immunity and stromal cell development because of its abundant expression in these tissues.

another tissue or cell sample taken from an individual having such a disorder, relative to

the standard gene expression level, i.e., the expression level in healthy tissue or bodily

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# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 103

This gene is expressed primarily in infant brain and placenta.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of the brain and nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, especially brain, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and placenta, cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue

The tissue distribution indicates that the protein product of this clone is useful for detecting defects of the brain, especially in young children.

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or bodily fluid from an individual not having the disorder.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 105

This gene is expressed primarily in human osteoclastoma and to a lesser extent in human pancreas tumor.

another tissue or cell sample taken from an individual having such a disorder, relative to immunological probes for differential identification of the tissue(s) or cell type(s). For a expression of this gene at significantly higher or lower levels may be routinely detected the standard gene expression level, i.e., the expression level in healthy tissue or bodily in certain tissues and cell types (e.g., bone and pancreas, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or biological sample and for diagnosis of diseases and conditions which include, but are number of disorders of the above tissues or cells, particularly in transformed tissues, Therefore, polynucleotides and polypeptides of the invention are useful as polypeptides and antibodies directed to these polypeptides are useful in providing not limited to, cancer particularly osteoclastoma and pancreatic tumor. Similarly, reagents for differential identification of the tissue(s) or cell type(s) present in a fluid from an individual not having the disorder. 22 ಜ 2

The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of some types of tumors, particularly pancreatic cancer and osteoclastoma.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 106

This gene is expressed primarily in fetal liver/spleen, and to a lesser extent in rated T-Cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the

- immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver, spleen, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the
- 5 expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of immune disorders.

# 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 107

This gene is expressed primarily in human embryo and to a lesser extent in spleen and chronic lymphocytic leukemia.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a hiological sample and for diagnosis of diseases and conditions; leukemia. Similarly,

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biological sample and for diagnosis of diseases and conditions: leukemia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune or hemopoietic systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, spleen, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that the protein product of this clone is useful for the diagnosis and treatment of leukemia.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 108

This gene is expressed primarily in placenta, and to a lesser extent in early stage human brain and in lung.

- reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: fetal developmental abnormalities. Similarly, polypeptides and antibodies directed to these polypeptides useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly in fetal and amniotic tissue, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, brain and other tissue of the nervous system, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution indicates that the protein product of this is useful for production of growth factor(s) associated with fetal development. Preferred polypeptides comprise the full-length polypeptide shown in the sequence listing, truncated however, at the amino terminus and beginning with QTE.

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### FEATURES OF PROTEIN ENCODED BY GENE NO: 109

This gene is expressed primarily in fetal spleen, and to a lesser extent in B-Cell

25 lymphoma and T-Cell lymphoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., spleen and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder,

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or bodily fluid from an individual not having the disorder.

relative to the standard gene expression level, i.e., the expression level in healthy tissue

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The tissue distribution indicates that the protein product of this clone is useful for the treatment and diagnosis of human lymphomas

### FEATURES OF PROTEIN ENCODED BY GENE NO: 110

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The translation product of this gene shares sequence homology with sarcoma amplified sequence (SAS), a tetraspan receptor which is thought to be important in malignant fibrous histiocytoma and liposarcoma. This gene is expressed primarily in human osteoclastoma, and to a lesser extent in pineal gland and infant brain.

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polypeptides are useful in providing immunological probes for differential identification relative to the standard gene expression level, i.e., the expression level in healthy tissue nistiocytoma and liposarcoma. Similarly, polypeptides and antibodies directed to these wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, fluid) or another tissue or cell sample taken from an individual having such a disorder, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone, Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: malignant fibrous pineal gland, and brain and other tissue of the nervous system, and cancerous and reagents for differential identification of the tissue(s) or cell type(s) present in a or bodily fluid from an individual not having the disorder. 15

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indicate that the protein product of this clone is useful for treatment of, osteosarcoma, The tissue distribution and homology to sarcoma amplified sequence (SAS) malignant fibrous histiocytoma and liposarcoma and related cancers, particularly sarcomas.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 111

The translation product of this gene shares sequence homology with 6.8K

This gene is expressed primarily in Wilm's tumor and to a lesser extent in proteolipid protein, mitochondrial - bovine. 2

cerebellum and placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Wilm's tumor.

providing immunblogical probes for differential identification of the tissue(s) or cell

Similarly, polypeptides and antibodies directed to these polypeptides are useful in

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may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and placenta, and cancerous and wounded tissues) or bodily fluids immune or renal systems, expression of this gene at significantly higher or lower levels sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an type(s). For a number of disorders of the above tissues or cells, particularly of the (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell

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the protein product of this clone is useful for diagnostic and therapeutics associated with The tissue distribution and homology to 6.8K proteolipid protein indicate that tumors, particularly Wilm's tumor disease. 2

individual not having the disorder.

### FEATURES OF PROTEIN ENCODED BY GENE NO: 112

This gene is expressed primarily in embryonic tissue and to a lesser extent in

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biological sample and for diagnosis of diseases and conditions: immune disorders. osteoblasts, endothelial cells, macrophages (GM-CSF treated), and bone marrow. Therefore, polynucleotides and polypeptides of the invention are useful as Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a

endothelial cells, blood cells and bone marrow, and cancerous and wounded tissues) or immune system, expression of this gene at significantly higher or lower levels may be providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the routinely detected in certain tissues and cell types (e.g., embryonic tissue, bone, ន

tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another fluid from an individual not having the disorder. 22

corresponding to this gene are useful for treatment or diagnosis of immune disorders. Preferred polypeptides encoded by this gene comprise the following amino acid sequence: MITDVQLAIFANMLGVSLFLLVVLYHYVAVNNPKKQE (SEQ ID NO: 636). The tissue distribution indicates that polynucleotides and polypeptides ജ

## FEATURES OF PROTEIN ENCODED BY GENE NO: 113

This gene is expressed primarily in hepatocellular tumor, and to a lesser extent in fetal liver/spleen. 35

of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification particularly of the hepatic system, expression of this gene at significantly higher or biological sample and for diagnosis of diseases and conditions: tumors, particularly hepatocellular tumors. Similarly, polypeptides and antibodies directed to these reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

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5 urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, the expression level in healthy tissue or bodily fluid from an individual not having the individual having such a disorder, relative to the standard gene expression level, i.e.,

lower levels may be routinely detected in certain tissues and cell types (e.g., liver, and

for diagnosis and treatment of tumors, particularly hepatocellular tumors. The tissue distribution indicates that the protein product of this clone is useful

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 114

colleagues (Nature 389, 300-306 (1997)). In addition, the predicted translation product apoptosis. The sequence of this gene has since been published by Polyak and EI24 which is also thought to be important in p53 mediated apoptosis. of this contig exhibits very high sequence homology with a murine gene denoted as identity with the human Pig8 gene which is thought to be important in p53 mediated The translation product of this gene exhibits a very high degree of sequence

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lesser extent in bone marrow, fetal liver, and prostate. This gene is expressed primarily in infant brain and activated T-cells and to a

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reagents for differential identification of the tissue(s) or cell type(s) present in a not limited to, and tissue damage by radiation and anti-cancer drugs. Similarly, biological sample and for diagnosis of diseases and conditions which include, but are Therefore, polynucleotides and polypeptides of the invention are useful as

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ၓ polypeptides and antibodies directed to these polypeptides are useful in providing nervous system, blood cells, bone marrow, liver, and prostate, and cancerous and routinely detected in certain tissues and cell types (e.g., brain and other tissue of the number of disorders of the above tissues or cells, particularly of the nervous and immunological probes for differential identification of the tissue(s) or cell type(s). For a fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal immune systems, expression of this gene at significantly higher or lower levels may be

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or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue

also be useful in the treatment of hematopoietic disorders and in boosting numbers of is upregulated in cells undergoing such treatment where p53 was overexpressed. It may etoposide, hydroperoxycyclophosphamide, and X-irradiation, since this protein product preventing apoptosis in patients being treated with anti-oncogenic drugs such as indicate that polynucleotides and polypeptides corresponding to this gene are useful for hematopoietic stem cells by interfering with the apoptosis of progenitor cells. The The tissue distribution and homology to human Pig8 and murine E124 genes

5 5 mature polypeptide is predicted to comprise the following amino acid sequence: EPRIVSRIFQCCAWNGGVFWFSLLLFYRVFIPVLQSVTARIIGDPSLHGDVWSWLEFFLTSIFSA FPSPHPSPAKLKATAGH (SEQ ID NO: 637). Accordingly, polypeptides comprising the SSYIISGCLFSILFPLFIISANEAKTPGKAYLFQLRLFSLVVFLSNRLFHKTVYLQSALSSSTSAEK FPIHLVGQLVSLLHMSLLYSLYCFEYRWFNKGIEMHQRLSNIERNWPYYFGFGLPLAFLTAMQ LWYLPLFYLSKYVNAIWFQDIADLAFEYSGRKPHPFPSYSKIIADMLFNLLLQALFLIQGMFYSI eemadsvktflqdlargikdsiwgictiskldariqqkreeqrrrassvlaqrraqsierkqes foregoing amino acid sequence are provided as are polynucleotides encoded such

## FEATURES OF PROTEIN ENCODED BY GENE NO: 115

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multiple sclerosis. This gene is expressed primarily in stromal cells and to a lesser extent in

reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

25 30 biological sample and for diagnosis of diseases and conditions: affecting the nervous routinely detected in certain tissues and cell types (e.g., stromal cells and cancerous and in providing immunological probes for differential identification of the tissue(s) or cell system. Similarly, polypeptides and antibodies directed to these polypeptides are useful type(s). For a number of disorders of the above tissues or cells, particularly of the nervous system, expression of this gene at significantly higher or lower levels may be or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal

and other autoimmune diseases. corresponding to this gene are useful for treatment and diagnosis of multiple sclerosis The tissue distribution indicates that polynucleotides and polypeptides

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 116

This gene is expressed primarily in the gall bladder

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: gall stones or infection of the digestive system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

significantly of the digestive system or renal system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., gall bladder and tissue of the digestive system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for possible prevention of digestive disorders where there may be a lack of digestive enzymes produced or in the detection and possible prevention of gall stones.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 117

The translation product of this gene shares sequence homology with dystrophin gene which is thought to be important in building and maintenance of muscles. This gene is expressed primarily in placenta and to a lesser extent in fetal brain

This gene is expressed primarily in placenta and to a lesser extent in fetal brain and fetal liver, and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as

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reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: muscular dystropy, Duchenne and Becker's muscular dystropies. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the skeletal muscle system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, brain and other tissue of the nervous system, muscle,

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an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution and homology to the dystrophin gene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diseases related the degenerative myopathies that are characterized by the weakness and atrophy of muscles without neural degradation; such as Duchenne and Becker's muscular dystropies.

# 10 FEATURES OF PROTEIN ENCODED BY GENE NO: 118

This gene is expressed primarily in olfactory tissue and to a lesser extent in

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: connective ussue diseases; chondrosarcoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

particularly of the connective tissue, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., olfactory tissue and cartilage, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for tumors of connective tissues, osteoarthritis and the treatment and diagnosis of chondrosarcoma.

# 30 FEATURES OF PROTEIN ENCODED BY GENE NO: 119

This gene is expressed primarily in Activated Neutrophils and to a lesser extent in fetal spleen, and CD34 positive cells from cord blood.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: allergies, defects in hematopoiesis and inflammation. Similarly, polypeptides and antibodies directed to

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plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from

liver, and spicen, and cancerous and wounded tissues) or bodily fluids (e.g., serum,

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these polypeptides are useful in providing immunological probes for differential

or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal in certain tissues and cell types (e.g., blood cells, and spleen, and cancerous and expression of this gene at significantly higher or lower levels may be routinely detected tissues or cells, particularly of the immune system and hematopoiesis system the identification of the tissue(s) or cell type(s). For a number of disorders of the above fluid) or another tissue or cell sample taken from an individual having such a disorder,

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corresponding to this gene are useful for reducing the allergic effects felt by allergy suffers by neutralizing the activity of the immune system, especially since neutrophils are abundant in persons suffering from allergies and other inflammatory conditions The tissue distribution indicates that polynucleotides and polypeptides

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 120

2 binding protein  $\Pi$  which is thought to be important in RNA binding for transcription of The translation product of this gene shares sequence homology with poly A

This gene is expressed primarily in colon and to a lesser extent in brain and

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25 မ number of disorders of the above tissues or cells, particularly of the immune and polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: colon cancer. Similarly, reagents for differential identification of the tissue(s) or cell type(s) present in a routinely detected in certain tissues and cell types (e.g., colon, tissue and cells of the digestive system, expression of this gene at significantly higher or lower levels may be or bodily fluid from an individual not having the disorder. immune system, and brain or other tissue of the nervous system, and cancerous and relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal immunological probes for differential identification of the tissue(s) or cell type(s). For a Therefore, polynucleotides and polypeptides of the invention are useful as

and treatment of colon cancer and other disorders of the digestive system. polynucleotides and polypeptides corresponding to this gene are useful for detection The tissue distribution and homology to poly A binding protein II indicate that

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 121

diphosphoglucose 4.6 dehydrase which is thought to be important in the metabolism of The translation product of this gene shares sequence homology with thymidine

This gene is expressed primarily in fetal liver and spleen and to a lesser extent in

polypeptides and antibodies directed to these polypeptides are useful in providing reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: diabetes. Similarly, Therefore, polynucleotides and polypeptides of the invention are useful as

- 5 5 nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, the expression level in healthy tissue or bodily fluid from an individual not having the an individual having such a disorder, relative to the standard gene expression level, i.e., plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from in certain tissues and cell types (e.g., liver, spleen, and brain and other tissue of the expression of this gene at significantly higher or lower levels may be routinely detected number of disorders of the above tissues or cells, particularly of the endocrine system, immunological probes for differential identification of the tissue(s) or cell type(s). For a
- 20 dehydrase indicate that polynucleotides and polypeptides corresponding to this gene are in the metabolism of sugar in to its more basic components. useful for treatment of persons with diabetes since it appears that this protein is needed The tissue distribution and homology to thymidine diphospoglucose 4.6

#### 23 FEATURES OF PROTEIN ENCODED BY GENE NO: 122

and VIII. Defects in the circulating levels of ceruloplasmin (aceruloplasminemia) have and copper. Ceruloplasmin also contains domains with homology to clotting factors V ceruloplasmin which is thought to be important in the metabolism and transport of iron The translation product of this gene shares sequence homology with

been associated with certain disease conditions such as Wilson disease, and the accompanying hepatolenticular degeneration.

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endothelial cells This gene is expressed primarily in brain and retina and to a lesser extent in

ઝ reagents for differential identification of the tissue(s) or cell type(s) present in a defects in iron metabolism; aceruloplasminemia not characterized by defects in the biological sample and for diagnosis of diseases and conditions: diseases marked by Therefore, polynucleotides and polypeptides of the invention are useful as

known ceruloplasmin gene locus; nonclassical Wilson disease; movement disorders; and tumors derived from a brain tissue origin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of

the above tissues or cells, particularly of the brain, retina, and nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, retinal tissue, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ceruloplasmin indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment of patients with aceruloplasminemia, or other defects in iron and/or copper metabolism. Mutations in this locus could also be diagnostic for patients currently experiencing or predicted to experience aceruloplasminemia.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 123

This gene is expressed primarily in brain and B cell lymphoma and to a lesser extent in fetal liver and spleen.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: B cell lymphoma; tumors and diseases of the brain and/or spleen; hematopoietic defects. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and hematopoietic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, blood cells, liver, and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides , corresponding to this gene are useful for treatment of disorders in neuronal,

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hematopoietic, and immune systems. It could potentially be useful for neurodegenerative disorders and neuronal and/or hematopoietic cell survival or proliferation.

## 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 124

This gene is expressed primarily in osteoclastoma, dermatofibrosarcoma, and B cell lymphoma and to a lesser extent in endothelial cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer in particular osteoclastoma, dermatofibrosarcoma, and B cell lymphoma. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of

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1.5 the bone, immune, and circulatory system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., bone, epidermis, blood cells, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of cancers and lymphoma; osteoporosis, and the control of cell proliferation and/or differentiation.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 125

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This gene is expressed primarily in immune tissues and hematopoietic cells, particularly in activated T cells and neutrophils, spleen, and fetal liver, and to a lesser extent in infant adrenal gland.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects in T cell activation; hematopoietic disorders; tumors of a hematopoietic and/or adrenal gland origin. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the hematopoietic and/or endocrine systems, expression of this gene at significantly higher

or lower levels may be routinely detected in certain tissues and cell types (e.g., cells and tissues of the immune system, hematopoietic cells, blood cells, liver, and adrenal gland, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for immune and/or hematopoietic disorders; diseases related to proliferation and/or differentiation of hematopoietic cells; defects in T cell and neutrophil activation and responsiveness; and endocrine and/or metabolic disorders, particularly of early childhood.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 126

This gene is expressed primarily in placenta and endothelial cells and to a lesser extent in melanocytes and embryonic tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors of an endothelial cell origin; angiogenesis associated with tumor development and metastasis. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the vascular system and developing embryo, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, endothelial

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may be routinely detected in certain tissues and cell types (e.g., placenta, endothelial cells, melanocytes, and embryonic tissues, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of developmental disorders; inhibition of angiogenesis; and vascular patterning.

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# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 127

This gene is expressed primarily in endothelial cells and hematopoietic tissues, including spleen, tonsils, leukocytes, and both B- and T-cell lymphomas.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors of an endothelial cell and/or hematopoietic origin; leukemias and lymphomas. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and vascular systems expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., endothelial cells, hematopoietic cells, spleen,

tonsils, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the manipulation of angiogenesis; the differentiation and morphogenesis of endothelial cells; the proliferation and/or differentiation of hematopoietic cells; and the commitment of hematopoietic cells to distinct cell lineages.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 128

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This gene is expressed primarily in kidney medulla and to a lesser extent in spleen from chronic myelogenous leukemia patients, prostate cancer, and some other tissues.

25 30 35 reagents for differential identification of the tissue(s) or cell type(s) present in a for differential identification of the tissue(s) or cell type(s). For a number of disorders biological sample and for diagnosis of diseases and conditions: tumors of a kidney of the above tissues or cells, particularly of the kidney and spleen, expression of this origin; chromic myelogenous leukemia; prostate cancer. Similarly, polypeptides and standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another and cell types (e.g., kidney, spleen, and prostate, and cancerous and wounded tissues) gene at significantly higher or lower levels may be routinely detected in certain tissues antibodies directed to these polypeptides are useful in providing immunological probes fluid from an individual not having the disorder. Therefore, polynucleotides and polypeptides of the invention are useful as

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It may also be useful for the enhancement of kidney tubule regeneration in the treatment disorders and cancer, particularly chromic myelogenous leukemia and prostate cancer. corresponding to this gene are useful for the diagnosis and treatment of kidney The tissue distribution indicates that polynucleotides and polypeptides of acute renal failure.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 129

This gene is expressed primarily in adult and infant brain and to a lesser extent in mesenchymal or fibroblast cells, as well as tissues with a mesenchymal origin.

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biological sample and for diagnosis of diseases and conditions: tumors of a brain and/or immunological probes for differential identification of the tissue(s) or cell type(s). For a mesenchymal cells and tissues, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell Therefore, polynucleotides and polypeptides of the invention are useful as tissue of the nervous system and cancerous and wounded tissues) or bodily fluids polypeptides and antibodies directed to these polypeptides are useful in providing number of disorders of the above tissues or cells, particularly of the brain and of reagents for differential identification of the tissue(s) or cell type(s) present in a mesenchymal origin; neurodegenerative disorders; cancer; fibrosis. Similarly,

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mesenchymal origin; neurodegenerative disorders; cancer; and fibrosis, based upon the expression of this gene within those tissues. Fibrosis is considered as mesenchymal expression level, i.e., the expression level in healthy tissue or bodily fluid from an corresponding to this gene are useful for the diagnosis of tumors of a brain and/or cells and fibroblasts are the primary cellular targets involved in this pathological The tissue distribution indicates that polynucleotides and polypeptides individual not having the disorder. 25

sample taken from an individual having such a disorder, relative to the standard gene

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condition.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 130

This gene is expressed primarily in hepatocellular cancer and to a lesser extent in fetal tissues as well as testes tumor.

biological sample and for diagnosis of diseases and conditions: liver cancer. Similarly, Therefore, polynucleotides and polypeptides of the invention are useful as polypeptides and ahtibodies directed to these polypeptides are useful in providing reagents for differential identification of the tissue(s) or cell type(s) present in a

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immunological probes for differential identification of the tissue(s) or cell type(s). For a expression of this gene at significantly higher or lower levels may be routinely detected number of disorders of the above tissues or cells, particularly of the digestive system, in certain tissues and cell types (e.g., liver, fetal tissue, and testes and other

- reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., he expression level in healthy tissue or bodily fluid from an individual not having the S
- corresponding to this gene are useful for diagnosis and treatment of liver cancer. The tissue distribution indicates that polynucleotides and polypeptides 2

## FEATURES OF PROTEIN ENCODED BY GENE NO: 131

This gene is expressed only in infant early brain.

- diseases of the nervous system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, higher or lower levels may be routinely detected in certain tissues and cell lypes (e.g., brain and other tissue of the nervous system and cancerous and wounded tissues) or particularly of the brain and nervous system, expression of this gene at significantly bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: development and reagents for differential identification of the tissue(s) or cell type(s) present in a 2 2
  - tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder. 25

corresponding to this gene are useful for treating diseases of the brain in children and in treating nervous system disorders such as Alzheimer's disease, schizophrenia, The tissue distribution indicates that polynucleotides and polypeptides dementia, depression, etc.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 132

This gene is expressed primarily in brain and to a lesser extent in glioblastoma. biological sample and for diagnosis of diseases and conditions: Alzheimer's disease, Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

schizophrenia, depression, mania, and dementia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain and nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating brain disorders such as Alzheimer's disease, schizophrenia, depression, mania, and dementia.

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healthy tissue or bodily fluid from an individual not having the disorder

# 15 FEATURES OF PROTEIN ENCODED BY GENE NO: 133

The translation product of this gene shares sequence homology with ribitol dehydrogenase of bacteria which is thought to be important in metabolism of sugars.

This gene is expressed primarily in macrophage and to a lesser extent in T-cell

lymphoma and lung.

reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tissue destruction in inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ribitol dehydrogenase indicate that polynucleotides and polypeptides corresponding to this gene are useful for altering macrophage metabolism in diseases such as inflammation where macrophages are causing excess tissue destruction.

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#### FEATURES OF PROTEIN ENCODED BY GENE NO: 1:

This gene is expressed primarily in pancreatic tumor and to a lesser extent in synovial sarcoma.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to,. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the endocrine and connective tissue systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., pancreas, and synovial tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating and diagnosing various cancers.

# 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 135

This gene is expressed primarily in T cell lines such as Raji and to a lesser extent in infant brain.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune system disorders and inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or

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30 lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating and diagnosing inflammatory diseases

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such as rheumatoid arthritis, sepsis, inflammatory bowel disease, and psoriasis, as well as neutropenia.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 136

The translation product of this gene shares high sequence homology with SAR1 subfamily of GTP-binding proteins which is thought to be important in vesicular transport in mammalian cells.

This gene is expressed primarily in serum-stimulated smooth muscle cells and to a lesser extent in a T-cell lymphoma.

polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, cells, and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., particularly of the muscular system, expression of this gene at significantly higher or serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample lower levels may be routinely detected in certain tissues and cell types (e.g., blood expression level, i.e., the expression level in healthy tissue or bodily fluid from an Therefore, polynucleotides and polypeptides of the invention are useful as biological sample and for diagnosis of diseases and conditions: diseases affecting reagents for differential identification of the tissue(s) or cell type(s) present in a taken from an individual having such a disorder, relative to the standard gene vesicular transport. Similarly, polypeptides and antibodies directed to these individual not having the disorder. 2 15 8

polynucleotides and polypeptides corresponding to this gene are useful for gene therapy in treating the large number of diseases involved in defective vesicular transport within

## FEATURES OF PROTEIN ENCODED BY GENE NO: 137

The translation product of this gene shares sequence homology with a protein

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found in *C. elegans* cosmid F25B5.

This gene is expressed primarily in a fetal tissues and to a lesser extent in melanocytes.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: abnormal fetal development, especially of the pulmonary system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes

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for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the fetal pulmonary system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., fetal tissue, pulmonary tissue, and melanocytes, and

5 cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides

10 corresponding to this gene are useful for treatment and diagnosis of diseases affecting
the pulmonary system, such as emphysema.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 138

This gene is expressed primarily in gall bladder and to a lesser extent in smooth

muscle.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: digestive system disease and gall bladder problems. Similarly, polypeptides and antibodies directed to these

- polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the digestive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., gall bladder and tissue of the digestive system, and smooth muscle, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal
  - wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treating diseases of the digestive system.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 139

This gene is expressed primarily in placenta and to a lesser extent in brain.

Therefore, polynucleotides and polypeptides of the invention are useful as
reagents for differential identification of the tissue(s) or cell type(s) present in a
biological sample and for diagnosis of diseases and conditions: abnormal fetal
development. Similarly, polypeptides and antibodies directed to these polypeptides are

expression level, i.e., the expression level in healthy tissue or bodily fluid from an sample taken from an individual having such a disorder, relative to the standard gene (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids or cell type(s). For a number of disorders of the above tissues or cells, particularly of individual not having the disorder. be routinely detected in certain tissues and cell types (e.g., placenta, and brain and other developing tissues, expression of this gene at significantly higher or lower levels may useful in providing immunological probes for differential identification of the tissue(s

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5 corresponding to this gene are useful for treating and diagnosing abnormal fetal The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 140

2 ovary, prostate cancer, and activated monocytes. This gene is expressed primarily in smooth muscle and to a lesser extent in

biological sample and for diagnosis of diseases and conditions: hypertension and reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

23 20 atherosclerosis. Similarly, polypeptides and antibodies directed to these polypeptides muscle, ovary and other reproductive tissue, prostate, and blood cells, and cancerous are useful in providing immunological probes for differential identification of the lower levels may be routinely detected in certain tissues and cell types (e.g., smooth particularly of the circulatory system, expression of this gene at significantly higher or tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

healthy tissue or bodily fluid from an individual not having the disorder. disorder, relative to the standard gene expression level, i.e., the expression level in spinal fluid) or another tissue or cell sample taken from an individual having such a and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or

such as hypertension, atherosclerosis, etc. corresponding to this gene are useful for treating diseases of the circulatory system The tissue distribution indicates that polynucleotides and polypeptides

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 141

35 and bone marrow This gene is expressed primarily in fetal spleen and to a lesser extent in placenta

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of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification significantly higher or lower levels may be routinely detected in certain tissues and cell particularly of the circulatory and pulmonary systems, expression of this gene at diseases affecting blood cells. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: anemia and other reagents for differential identification of the tissue(s) or cell type(s) present in a types (e.g., spleen, placenta, bone marrow, and blood cells, and cancerous and Therefore, polynucleotides and polypeptides of the invention are useful as

5 or bodily fluid from an individual not having the disorder relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal

5 corresponding to this gene are useful for the generation of red and white blood cells and for the diagnosis of disease of these cells. The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 142

8 murine tetracycline/sugar transporter molecule recently reported by Matsuo and colleagues (Biochem. Biophys. Res. Commun. 238 (1), 126-129 (1997)). The predicted translation product of this contig is a human homolog of the

This gene is expressed primarily in synovium and to a lesser extent in

reagents for differential identification of the tissue(s) or cell type(s) present in a or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) inflammation. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: rheumatoid arthritis and Therefore, polynucleotides and polypeptides of the invention are useful as

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ဗ 33expression level, i.e., the expression level in healthy tissue or bodily fluid from an the immune and lymphatic systems, expression of this gene at significantly higher or serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample tissue, and endothelial cells, and cancerous and wounded tissues) or bodily fluids (e.g. lower levels may be routinely detected in certain tissues and cell types (e.g., synovial taken from an individual having such a disorder, relative to the standard gene

individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment and diagnosis of inflammatory diseases, such as rheumatoid arthritis, leukemia, neutropenia, inflammatory bowel disease, psoriasis, sepsis, and the like.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 143

This gene is expressed primarily in placenta and to a lesser extent in melanocyte, retal liver and spleen, and bone marrow.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: abnormal early development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, melanocytes, liver, spleen, and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and diagnosis of abnormal early development phenomena and diseases.

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# 25 FEATURES OF PROTEIN ENCODED BY GENE NO: 144

This gene is expressed primarily in fetal liver and spleen.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: anemia and neutropenia. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and blood systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., liver and spleen, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine,

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synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual

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having such a disorder, relative to the standard gene expression level, i.e., the

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expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful in hematopoeisis and bone marrow regeneration as it is most abundant in fetal tissues responsible for the generation of hematopoeitic

## FEATURES OF PROTEIN ENCODED BY GENE NO: 145

The translation product of this gene shares sequence homology with protein tyrosine phosphatase which is thought to be important in transducing signal to activate cells such as T cell, B cell and other cell types.

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This gene is expressed primarily in T cells and tissues in early stages of development and to a lesser extent in cancers.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immuno-related diseases and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

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20 particularly of the immune system expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic and fetal tissue, undifferentiated cells, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily

fluid from an individual not having the disorder.

The tissue distribution and homology to the protein tyrosine phosphatase family indicate that polynucleotides and polypeptides corresponding to this gene are useful for modulating the immune system.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 146

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This gene is expressed primarily in T cell and to a lesser extent in B cell, macrophages and tumor tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immuno-disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in

wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal or bodily fluid from an individual not having the disorder relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder, routinely detected in certain tissues and cell types (e.g., blood cells, and cancerous and immune system, expression of this gene at significantly higher or lower levels may be type(s). For a number of disorders of the above tissues or cells, particularly of the providing immunological probes for differential identification of the tissue(s) or cel

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5 corresponding to this gene are useful for regulating the immune system therefore can be used in treating diseases such as autoimmune diseases and cancers The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 147

cells, testis tumor, ovarian cancer, uterine cancer. This gene is expressed primarily in placenta and to a lesser extent in endothelial

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not limited to cancer. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions which include, but are reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

8 of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, polypeptides are useful in providing immunological probes for differential identification endothelial cells, testis and ovary and other reproductive tissue, and cancerous and particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta,

25 wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal or bodily fluid from an individual not having the disorder. relative to the standard gene expression level, i.e., the expression level in healthy tissue fluid) or another tissue or cell sample taken from an individual having such a disorder,

corresponding to this gene are useful for diagnosis and treatment of cancers The tissue distribution indicates that polynucleotides and polypeptides

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 148

Genet. 17, 40-48 (1997).) group cloned the human Torsin A gene. (See, Accession No. 2358279; see also Nature This sequence has significant homology to mouse torsin A. Recently, another

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standard gene expression level, i.e., the expression level in healthy tissue or bodily tissue or cell sample taken from an individual having such a disorder, relative to the

fluid from an individual not having the disorder.

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lesser extent in fetal lung, fetal liver, fetal brain, adult brain and tumor tissues This gene is expressed primarily in osteoclastoma, T-cell, and placenta and to a

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5 of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, cells, bone, placenta, lung, liver, and brain and other tissues of the nervous system, particularly of the hematopoiesis system, expression of this gene at significantly higher polypeptides are useful in providing immunological probes for differential identification hematopoiesis and cancers. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: disease conditions in expression level in healthy tissue or bodily fluid from an individual not having the having such a disorder, relative to the standard gene expression level, i.e., the synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual or lower levels may be routinely detected in certain tissues and cell types (e.g., blood reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

corresponding to this gene are useful for treating blood related diseases such as deficiencies in red blood cell, white blood cell, platelet and other hematopoiesis cells The tissue distribution indicates that polynucleotides and polypeptides

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 149

20 spleen and osteoclastoma. endothelial cells and to a lesser extent in monocyte, dendritic cell, bone marrow, salivary gland, colon cancer, stomach cancer, pancreatic tumor, uterine cancer, fetal This gene is expressed primarily in T cell, prostate and prostate cancer,

25 မွ reagents for differential identification of the tissue(s) or cell type(s) present in a particularly of the immune system, expression of this gene at significantly higher or of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, diseases and cancers. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: immuno-related stomach, pancreas, uterus, spleen and bone, and cancerous and wounded tissues) or cells, prostate, endothelial cells, dendritic cells, bone marrow, salivary gland, colon, polypeptides are useful in providing immunological probes for differential identification bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another lower levels may be routinely detected in certain tissues and cell types (e.g., blood Therefore, polynucleotides and polypeptides of the invention are useful as

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for treatment of cancers.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 150

This gene was recently cloned by another group, calling it eIF3-p66. (See Accession No. 2351378.) This gene plays a role in RNA binding and macromolecular assembly, and therefore, any mutations in this gene would likely result in a diseased phenotype. Preferred polypeptide fragments comprise the amino acid sequence: MAKFMTPVIQDNPSGWGPCAVPEQFRDMPYQPFSKGDRLGRVADWTGATYQDKRYTNKYSS

10 QFGGGSQYAYFHEEDESSFQLVDTARTQKTAYQRNRMRFAQRNILRRDKDRRNMLQFNLQILP
KSAKQKERERIRLQKKFQKQFGVRQKWDQKSQKPRDSSVEVRSDWEVKEEMDFPQLMKMRY
LEVSEPQDIECCGALEYYDKAFDRITTRSEKPLRXXKRIFHTYTTTDDPVIRKLAKTQGNVFATD
AILATLMSCTRSVYSWDIVVQRVGSKLFPDKRDNSDFDLLTVSETANEPPQDEGNSFNSRML
AMEATYINHNFSQQCLRMGKERYNFPNPNPFVEDDMDKNEIASVAYRYRSGKLGDDIDLIVRC
15 EHDGVMTGANGEVSFINIKTLNEWDSRHCNGVDWRQKLDSQRGAVIATELKNNSYKLARWTC
CALLAGSEYLKLGYVSRYHVKDSSRHVILGTQQFKRNEFASQINLSVENAWGILRCVIDICMKL
EEGKYLILKDPNKQVIRVYSLPDGTFSS (SEQ ID NO: 638), as well as N-terminal and C-terminal deletions of this polypeptide fragment.

This gene is expressed primarily in T cell, bone marrow, embryo and endothelial cells and to a lesser extent in testis tumor and endometrial tumor.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: immune diseases and tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system and reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for immune disorders and cancers.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 151

This gene is expressed primarily in testis and to a lesser extent in T cell, spinal cord, placenta, neutrophil and monocyte.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: male reproductive and endocrine disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells,

particularly of the reproductive, immune and endocrine systems, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., testis and other reproductive tissue, blood cells, tissue of the nervous system, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for regulating immune and reproductive functions.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 152

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The translation product of this gene shares sequence homology with tyrosyl-tRNA synthetase which is thought to be important in cell growth.

This gene is expressed primarily in brain, liver, keratinocytes, tonsils, and

25 heart.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer autoimmune diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, liver, keratinocytes, tonsils, heart expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissues of the nervous system,

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35 liver, keratinocytes, tonsils and heart, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard

gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to tyrosyl-tRNA synthetase indicate that polynucleotides and polypeptides corresponding to this gene are useful for modulating cell growth.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 153

This gene is homologous to the Drosophila transcriptional regulator dre4. (See Accession No. 2511745.) Dre4 is a gene required for steroidogenesis in Drosophila

transcriptional regulator CDC68. Preferred polypeptide fragments comprise the amino acid sequence: KKRHTDVQFYTEVGEITTDLGKHQHMHDRDDLYAEQMEREMRHKLKTAFKN FIEKVEALTKEELEFEVPFRDLGFNGAPYRSTCLLQPTSSALVNATEWPPFVVTLDEVELIHFXR VQFHLKNFDMVIVYKDYSKKVTMINAIPVASLDPIKEWLNSCDLKYTEGVQSLNWTKIMKTIVD DPEGFFEQGGWSFL (SEQ ID NO: 639), as well as N-terminal and C-terminal deletions of this fragments. Also preferred are polynucleotide fragments encoding this polypeptide fragment.

This gene is expressed primarily in fetal liver, spleen, placenta, lung, T cell, yroid, testes.

25 20 છ or cell type(s). For a number of disorders of the above tissues or cells, particularly of plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from types (e.g., liver, spleen, placenta, lung, blood cells, thyroid, and testes and other significantly higher or lower levels may be routinely detected in certain tissues and cell useful in providing immunological probes for differential identification of the tissue(s) liver diseases. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: brain tumor, heart and reagents for differential identification of the tissue(s) or cell type(s) present in a an individual having such a disorder, relative to the standard gene expression level, i.e., reproductive tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, the fetal liver, spleen, placenta, lung, T cell, thyroid, testes expression of this gene at the expression level in healthy tissue or bodily fluid from an individual not having the Therefore, polynucleotides and polypeptides of the invention are useful as

# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 154

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This gene is expressed primarily in brain and to a lesser extent in fetal heart testis, spleen, lung.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: heart, liver and spleen diseases, immunological diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, fetal heart, testis, spleen, lung expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, heart, testes and other reproductive tissue, spleen, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 155

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Activation of T cells through the T cell antigen receptor (TCR) results in the rapid tyrosine phosphorylation of a number of cellular proteins, one of the earliest being a 100 kDa protein. This gene is the human equivalent of murine valosin containing protein (VCP). VCP is a member of a family of ATP binding, homo-oligomeric proteins, and the mammalian homolog of Saccharomyces cerevisiae cdc48p, a protein essential to the completion of mitosis in yeast. Both endogenous and expressed murine VCP are tyrosine phosphorylated in response to T cell activation. Thus we have identified a novel component of the TCR mediated tyrosine kinase activation pathway that may provide a link between TCR activation and cell cycle control.

This gene is expressed primarily in brain, liver, spleen, placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancer immunological disorders. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the brain, liver, spleen, placenta expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, liver, spleen, and placenta, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from

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an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to VCR indicate that polynucleotides and polypeptides corresponding to this gene are useful for treating cancer.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 156

response protein which is thought to be important in cell growth. A group recently cloned the human homolog of this gene, calling it insulin induced protein 1. (See The translation product of this gene shares sequence homology with rat growth

LVYNGVYQYTSPDFLYIRSWLPCIFFSGGVTVGNIGRQLAMGVPEKPHSD (SEQ ID NO: 640). polypeptide fragments comprise the amino acid sequence: RSGLGLGTIAFLATLITQF Accession No. 2358269, see also, Genomics 43 (3), 278-284 (1997).) Preferred as well as N-terminal and C-terminal deletions of this polypeptide fragment. Also preferred are polynucleotide fragments encoding these polypeptide fragments. 9 2

This gene is expressed primarily in brain, liver, placenta, heart, spleen,

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

not limited to, cancer immunological disorders. Similarly, polypeptides and antibodies biological sample and for diagnosis of diseases and conditions which include, but are differential identification of the tissue(s) or cell type(s). For a number of disorders of directed to these polypeptides are useful in providing immunological probes for 2

liver, placenta, heart, spleen, and lymphoid tissue, and cancerous and wounded tissues) expression of this gene at significantly higher or lower levels may be routinely detected or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily in certain tissues and cell types (e.g., brain and other tissue of the nervous system, the above tissues or cells, particularly of the brain, liver, placenta, heart, spleen. 8 25

polynucleotides and polypeptides corresponding to this gene are useful for modulating The tissue distribution and homology to growth-response protein indicate that cell growth.

fluid from an individual not having the disorder.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 157

This gene is expressed primarily in Glioblastoma, endometrial tumor, lymphoma and pancreas tumor.

lypes (e.g., endometrium, lymphoid tissue, pancreas, and tissue of the nervous system, antibodies directed to these polypeptides are useful in providing immunological probes significantly higher or lower levels may be routinely detected in certain tissues and cell for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at Therefore, polynucleotides and polypeptides of the invention are useful as and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, Endometrial tumor, lymphoma and pancreas tumor. Similarly, polypeptides and reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Glioblastoma, Ś 9

## FEATURES OF PROTEIN ENCODED BY GENE NO: 158

synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual

expression level in healthy tissue or bodily fluid from an individual not having the

disorder.

having such a disorder, relative to the standard gene expression level, i.e., t

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The translation product of this gene shares sequence homology with IGE receptor which is thought to be important in allergy and asthma.

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This gene is expressed primarily in T cell, and fetal liver.

polypeptides are useful in providing immunological probes for differential identification other immunological disorders. Similarly, polypeptides and antibodies directed to these biological sample and for diagnosis of diseases and conditions: allergy and asthma and of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune, expression of this gene at significantly higher or lower Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 23

liver, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, levels may be routinely detected in certain tissues and cell types (e.g., blood cells, and synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual expression level in healthy tissue or bodily fluid from an individual not having the having such a disorder, relative to the standard gene expression level, i.e., the ഉ 35

polynucleotides and polypeptides corresponding to this gene are useful for allergy and The tissue distribution and homology to IgE receptor indicate that

#### S FEATURES OF PROTEIN ENCODED BY GENE NO: 159

immunoglobin heavy chain which is thought to be important in immune response to the The translation product of this gene shares sequence homology with

activated T cell, monocyte and heart. This gene is expressed primarily in activated neutrophil and to a lesser extent in

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20 5 useful in providing immunological probes for differential identification of the tissue(s) such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial routinely detected in certain tissues and cell types (e.g., blood cells, and heart, and the immune, expression of this gene at significantly higher or lower levels may be or cell type(s). For a number of disorders of the above tissues or cells, particularly of and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: infection, inflammation reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful

useful for making the ligand to block specific antigen which cause certain disease. region indicate that polynucleotides and polypeptides corresponding to this gene are The tissue distribution and homology to immunoglobin heavy chain variable

in healthy tissue or bodily fluid from an individual not having the disorder.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 160

inactivation. inactive specific transcript protein which is thought to be important in X chromosome The translation product of this gene shares sequence homology with mouse X

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ovary tissue, ovarian cancer, frontal cortex and brain. This gene is expressed primarily in HSA 172 cell and to a lesser extent in normal

reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

3biological sample and for diagnosis of diseases and conditions: ovarian tumor, directed to these polypeptides are useful in providing immunological probes for schizophrenia and other neurological disorders. Similarly, polypeptides and antibodies

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sample taken from an individual having such a disorder, relative to the standard gene (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids the above tissues or cells, particularly of the immune and neural system, expression of differential identification of the tissue(s) or cell type(s). For a number of disorders of individual not having the disorder. expression level, i.e., the expression level in healthy tissue or bodily fluid from an tissues and cell types (e.g., ovary and other reproductive tissue, and brain and other this gene at significantly higher or lower levels may be routinely detected in certain

5 diagnosis and treatment of reproductive system tumors and CNS tumors. indicate that polynucleotides and polypeptides corresponding to this gene are useful for The tissue distribution and homology to X inactive specific transcript protein

#### FEATURES OF PROTEIN ENCODED BY GENE NO: 161

and prostate. This gene is expressed primarily in adipose cell and to a lesser extent in liver

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20 the adipose cell, expression of this gene at significantly higher or lower levels may be or cell type(s). For a number of disorders of the above tissues or cells, particularly of useful in providing immunological probes for differential identification of the tissue(s) disorder. Similarly, polypeptides and antibodies directed to these polypeptides are biological sample and for diagnosis of diseases and conditions: obesity and liver reagents for differential identification of the tissue(s) or cell type(s) present in a Therefore, polynucleotides and polypeptides of the invention are useful as

23 prostate, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma the expression level in healthy tissue or bodily fluid from an individual not having the individual having such a disorder, relative to the standard gene expression level, i.e., urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an

routinely detected in certain tissues and cell types (e.g., adipose cells, liver, and

ઝ corresponding to this gene are useful for treatment of obesity and liver disorder. The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 162

ઝ ubiquitin activating enzyme homolog which is thought to be important in protein posttraslation processing. The translation product of this gene shares sequence homology with yeast

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This gene is expressed primarily in stromal cell and to a lesser extent in retina,

H. Atrophic Endometrium, colon carcinoma and myeloid progenitor cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: defects of stromal cell development, neuronal growth disorders and tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., retinal cells, endometrium, colon, and bone marrow, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to ubiquitin-activating enzyme homolog indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of some type of tumors, fucosidosis and neuronal growth disorders.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 163

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This gene is expressed primarily in primary breast cancer and nemangiopericytoma and to a lesser extent in adult brain and cerebellum.

polypeptides are useful in providing immunological probes for differential identification such a disorder, relative to the standard gene expression level, i.e., the expression level biological sample and for diagnosis of diseases and conditions: breast cancer, leukemia cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial significantly higher or lower levels may be routinely detected in certain tissues and cell of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, fluid or spinal fluid) or another tissue or cell sample taken from an individual having Therefore, polynucleotides and polypeptides of the invention are useful as and cerebellum disorders. Similarly, polypeptides and antibodies directed to these particularly of the immune system and neural system, expression of this gene at types (e.g., mammary tissue, brain and other tissue of the nervous system, and reagents for differential identification of the tissue(s) or cell type(s) present in a n healthy tissue or bodily fluid from an individual not having the disorder. 22 റ്റ 35

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The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis or treatment of various tumors and disease involved in neural system.

# 5 FEATURES OF PROTEIN ENCODED BY GENE NO: 164

The translation product of this gene shares sequence homology with proline rich proteins. Recently, another group has also cloned this gene, calling it CD84 leukocyte antigen, a new member of the Ig superfamily. (See Accession No. U82988, see also, Blood 90 (6), 2398-2405 (1997).)

10 This gene is expressed primarily in Weizmann olfactory tissue and osteoclastoma and to a lesser extent in anergic T-cell.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: ostsis and immune disease. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., olfactory tissue, bone, and blood cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

The tissue distribution and homology to the Ig superfamily indicate that the protein product of this clone is useful for treatment of osteoporosis, autoimmune disease, and other immune disorders.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 165

30 This gene is expressed primarily in atrophic endometrium and colon cancer and to a lesser extent in some fetal tissues.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: tumors. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a

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number of disorders of the above tissues or cells, particularly of the immune system,

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in healthy tissue or bodily fluid from an individual not having the disorder. such a disorder, relative to the standard gene expression level, i.e., the expression level cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial in certain tissues and cell types (e.g., endometrium, colon, and fetal tissue, and expression of this gene at significantly higher or lower levels may be routinely detected fluid or spinal fluid) or another tissue or cell sample taken from an individual having

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corresponding to this gene are useful for diagnosis and treatment of tumors, specifically endometrium and colon tumors. The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 166

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the expression level in healthy tissue or bodily fluid from an individual not having the an individual having such a disorder, relative to the standard gene expression level, i.e.

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5 the amino acid sequence: VTQPKHLSASMGGSVEIPFSFYYPWELAXXPXVRISWRRGHFHG in activated monocyte. Although the predicted signal sequence is identified in Table 1, polynucleotide fragments encoding these polypeptide fragments. NO: 641), as well as N-terminal and C-terminal deletions. Also preferred are QSFYSTRPPSIHKDYVNRLFLNWTEGQESGFLRISNLRKEDQSVYFCRVELDTRRSG (SEQ ID other upstream sequences are also relevant. Preferred polypeptide fragments comprise This gene is expressed primarily in human primary breast cancer and to a lesser extent

20 reagents for differential identification of the tissue(s) or cell type(s) present in a number of disorders of the above tissues or cells, particularly of the immune system, polypeptides and antibodies directed to these polypeptides are useful in providing biological sample and for diagnosis of diseases and conditions: breast cancer. Similarly, immunological probes for differential identification of the tissue(s) or cell type(s). For a Therefore, polynucleotides and polypeptides of the invention are useful

30 expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., mammary tissue, and blood cells, and cancerous healthy tissue or bodily fluid from an individual not having the disorder spinal fluid) or another tissue or cell sample taken from an individual having such a and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or disorder, relative to the standard gene expression level, i.e., the expression level in

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corresponding to this gene are useful for diagnosis of breast cancer. The tissue distribution indicates that polynucleotides and polypeptides

## FEATURES OF PROTEIN ENCODED BY GENE NO: 167

35 used as a marker for linkage analysis for chromosome 9. lung. This gene has also been mapped to chromosomal location 9q34, and thus, can be This gene is expressed primarily in fetal tissues and to a lesser extent in adult

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tissues, and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, lower levels may be routinely detected in certain tissues and cell types (e.g., fetal particularly of the embryo tissues, expression of this gene at significantly higher or tissue(s) or,cell type(s). For a number of disorders of the above tissues or cells, are useful in providing immunological probes for differential identification of the biological sample. Similarly, polypeptides and antibodies directed to these polypeptides reagents for differential identification of the tissue(s) or cell type(s) present in a plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from Therefore, polynucleotides and polypeptides of the invention are useful as

## FEATURES OF PROTEIN ENCODED BY GENE NO: 168

5 Chain which is thought to be important in immune response. The translation product of this gene shares sequence homology with Ig Heavy

This gene is expressed primarily in prostate cancer tissue specifically

25 8 detected in certain tissues and cell types (e.g., prostate, tissue and cells of the immune biological sample and for diagnosis of diseases and conditions: prostate cancer. system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, prostate, expression of this gene at significantly higher or lower levels may be routinely providing immunological probes for differential identification of the tissue(s) or cell Similarly, polypeptides and antibodies directed to these polypeptides are useful in reagents for differential identification of the tissue(s) or cell type(s) present in a individual having such a disorder, relative to the standard gene expression level, i.e., type(s). For a number of disorders of the above tissues or cells, particularly of the urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an the expression level in healthy tissue or bodily fluid from an individual not having the Therefore, polynucleotides and polypeptides of the invention are useful as

## FEATURES OF PROTEIN ENCODED BY GENE NO: 169

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 $\frac{3}{2}$ acid metabolism. The gene represented by this contig has since been published by Hajra and colleagues (GenBank Accession No. U91316) acyl coenzyme-A hydrolase, which is thought to be important in neuron-specific fatty The translation product of this gene shares sequence homology with cytosolic

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This gene is expressed primarily in human pituitary gland and to a lesser extent in colorectal cancer tissue. This gene has also been observed in the LNCAP cell line.

and colon, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, the expression level in healthy tissue or bodily fluid from an individual not having the lower levels may be routinely detected in certain tissues and cell types (e.g., pituitary individual having such a disorder, relative to the standard gene expression level, i.e., familial and/or idiopathic origins. Similarly, polypeptides and antibodies directed to identification of the tissue(s) or cell type(s). For a number of disorders of the above ussues or cells, particularly blood, expression of this gene at significantly higher or biological sample and for diagnosis of diseases and conditions: hyperlipidemias of urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an Therefore, polynucleotides and polypeptides of the invention are useful as these polypeptides are useful in providing immunological probes for differential reagents for differential identification of the tissue(s) or cell type(s) present in a

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hydrolase indicate that polynucleotides and polypeptides corresponding to this gene are useful for the detection or treatment of hyperlipidemia disease states by virtue of the The tissue distribution and homology to rat cytosolic acyl coenzyme-A ability of specific drugs to activate the enzyme.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 170

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The translation product of this gene shares sequence homology with a Caenorhabditis elegans gene which is thought to be important in organism development.

This gene is expressed primarily in human synovial sarcoma tissue, bone marrow, and to a lesser extent in human brain. 53

antibodies directed to these polypeptides are useful in providing immunological probes routinely detected in certain tissues and cell types (e.g., synovial tissue, bone marrow, immune system, expression of this gene at significantly higher or lower levels may be for differential identification of the tissue(s) or cell type(s). For a number of disorders brain and other tissue of the nervous system, and cancerous and wounded tissues) or biological sample and for diagnosis of diseases and conditions which include, but are of the above tissues or cells, particularly of the bone, connective tissues and possibly bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another not limited to, of bone, specifically synovial sarcoma. Similarly, polypeptides and Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 35 2

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tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

polynucleotides and polypeptides corresponding to this gene are useful as a diagnostic The tissue distribution and homology to Caenorhabditis elegans indicate that and/or therapeutic modality directed at the detection and/or treatment of connective lissue sarcomas or other related bone diseases. S

## FEATURES OF PROTEIN ENCODED BY GENE NO: 171

6GlcNAc transferase which is thought to be important in the transfer and metabolism of invasiveness into the matrigel, and inhibition of cell attachment to collagen and laminin The translation product of this gene shares sequence homology with betalbeta1-6, N-acetylglucosamine. This gene product has previously been shown to suppress melanoma lung metastasis in both syngeneic and nude mice, decreased without affecting cell growth. 2 2

This gene is expressed primarily in human testes and prostate tissues, and to a lesser extent in kidney, medulla, and pancreas.

gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., testes and other reproductive tissue, prostate, kidney, pancreas, brain and biological sample and for diagnosis of diseases and conditions which include, but are differential identification of the tissue(s) or cell type(s). For a number of disorders of fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard not limited to, cancer particularly melanoma. Similarly, polypeptides and antibodies other tissue of the nervous system, and cancerous and wounded tissues) or bodily Therefore, polynucleotides and polypeptides of the invention are useful as directed to these polypeptides are useful in providing immunological probes for reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder. 2 ഉ

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The tissue distribution and homology to betal-6GlcNAc transferase indicate that therapeutic modalities directed at the detection and/or treatment of cancer, the metastasis of malignant tissue or cells. Defects in this potentially secreted enzyme may/play a role the protein product of this clone is useful for the development of diagnostic and/or in metastasis.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 172

This gene is expressed primarily in fetal spleen and liver.

5 5 S reagents for differential identification of the tissue(s) or cell type(s) present in a disorder, relative to the standard gene expression level, i.e., the expression level in and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or routinely detected in certain tissues and cell types (e.g., spleen and liver, and cancerous immune systems, expression of this gene at significantly higher or lower levels may be number of disorders of the above tissues or cells, particularly of the hematopoiesis and polypeptides and antibodies directed to these polypeptides are useful in providing Wilm's tumor disease, hepatic disorders, and hematopoietic disorders. Similarly, biological sample and for diagnosis of diseases and conditions: immune disorders healthy tissue or bodily fluid from an individual not having the disorder. spinal fluid) or another tissue or cell sample taken from an individual having such a immunological probes for differential identification of the tissue(s) or cell type(s). For Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution indicates that polynucleotides and polypeptides corresponding to this gene are useful for the treatment and identification of fetal defects along with correcting diseases that affect hematopoiesis and the immune system.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 173

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The translation product of this gene shares sequence homology with ret II oncogene which is thought to be important in Hirschsprung disease and many types of cancers.

This gene is expressed in multiple tissues including the lymphatic system, brain, and thyroid.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: Hirschsprung disease and multiple cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune and central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., lymphoid tissue, thyroid, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or

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another tissue or cell sample taken from an individual having such a disorder, relative to

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the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to ret II oncogene indicate that polynucleotides and polypeptides corresponding to this gene are useful for diagnosis and treatment of various cancers. It would also be useful for the diagnosis and treatment of Hirschsprung disease. Preferred polypeptides of the invention comprise the amino acid sequence: MEAQQVNEAESAREQLQXLHDQIAGQKASKQELETELERLKQEFHYIEEDLY RTKNTLQSRIKDRDEEIQKLRNQLTNKTLSNSSQSELENRLHQLTETLIQKQTMLESLSTEKNSL VFQLERLEQQMNSASGSSSNGSSINMSGIDNGEGTRLRNVPVLFNDTETNLAGMYGKVRKAAS OSDQFSIRLGIFLRRYPPARVFVIIYMALLHLWVMIVLLTYTPEM HHDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALLHLWVMIVLLTYTPEM HHDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALLHLWVMIVLTYTPEM HHDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALLHLUWMIVLTYTPEM HHDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALLHLUWMIVLTYTYTPEM HDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALLHLUWMIVLTYTYTPEM HDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALHLUWMIVLTYTYTPEM HDQPYGK (SEQ ID NO: 10 SIDQFSIRLGIFLRRYPPARVFVIIYMALHUM SIDQFTRLRWYMIVLTYTYTPEM SIDQFTRLRWYMIV SIDQFTRLRWYMIVLTYTYTPEM SIDQFTRLRWYMIV SIDQF

## FEATURES OF PROTEIN ENCODED BY GENE NO: 174

The translation product of this gene shares sequence homology with testis enhanced gene transcript which is thought to be important in regulation of human development.

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This gene is expressed primarily in infant brain and to a lesser extent in a variety of other tissues and cell types, including the prostate, testes, monocytes, macrophages, dendritic cells, keratinocytes, and adipocytes.

20 30 23 of the above tissues or cells, particularly of the brain and immune systems, expression cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial of this gene at significantly higher or lower levels may be routinely detected in certain for differential identification of the tissue(s) or cell type(s). For a number of disorders antibodies directed to these polypeptides are useful in providing immunological probes developmental, immune and inflammation disorders. Similarly, polypeptides and biological sample and for diagnosis of diseases and conditions: neurological, reagents for differential identification of the tissue(s) or cell type(s) present in a in healthy tissue or bodily fluid from an individual not having the disorder such a disorder, relative to the standard gene expression level, i.e., the expression level fluid or spinal fluid) or another tissue or cell sample taken from an individual having testes and other reproductive tissue, blood cells, keratinocytes, and adipocytes, and tissues and cell types (e.g., brain and other tissue of the nervous system, prostate, Therefore, polynucleotides and polypeptides of the invention are useful as

The tissue distribution and homology to testis enhanced gene transcript indicate
that the protein product of this clone is useful for diagnosis and treatment of disorders involving the developing brain and the immune system.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 175

This gene is expressed primarily in prostate and to a lesser extent in various other tissues, including placenta.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, cancers, especially of the prostate. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the prostate, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution indicates that the protein product of this clone is useful for diagnosis and treatment of prostate disorders and cancer. It may also be useful for the diagnosis and treatment of endocrine disorders.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 176

The translation product of this gene shares sequence homology with Sacchromyces cerevisiae YNT20 gene which is thought to be important in mitochondrial function.

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This gene is expressed at a particularly high level in muscle tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases related to such tissues and cell types including: muscle wasting diseases. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuromuscular system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., muscle and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e.,

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the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The tissue distribution and homology to the YNT20 gene indicate that this protein is useful for treatment and detection of neuromuscular diseases caused by loss of mitochondrial function. For example this gene or its protein product could be used in replacement therapy for such diseases.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 177

This gene is expressed primarily in the brain and to a lesser extentin kidney, placenta, smooth muscle, heart and lung.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: neuromuscular diseases, degenerative diseases of the central nervous system, and heart disease.

15 Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the neuromuscular system, central nervous system, and heart, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell

types (e.g., brain and other tissue of the nervous system, kidney, placenta, muscle, heart and lung, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

This gene or its protein product could also be used for replacement therapy for the above mentioned diseases.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 178

30 The translation product of this gene shares sequence homology with caldesmon which is thought to be important in the cellular response to changes in glucose levels. This gene is expressed primarily in multiple tissues including brain and retina.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions: central nervous system disorders and retinopathy. Similarly, polypeptides and antibodies directed to these polypeptides are

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useful in providing immunological probes for identification of the tissue(s) or cell

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type(s). For a number of disorders of the above tissues or cells, particularly of the CNS disorders and retinopathy, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, and retinal tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The tissue distribution and homology to caldesmon indicate that polynucleotides and polypeptides corresponding to this gene are useful for treatment of retinopathies.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 179

The translation product of this gene shares sequence homology with mouse fibrosin protein which is thought to be important in regulation of fibrinogenesis in

15 certain chronic inflammatory diseases.

This gene is expressed primarily in amniotic cells and breast tissue.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of breast cancer and abnormal embryo

- development. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., amniotic cells, and
- mammary tissue, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.
- The tissue distribution and homology to fibrosin indicate that the protein product of this clone is useful for treatment of breast cancer. This gene or its protein product could be used in replacement therapy for breast cancer. In addition the protein product of this gene is useful in the treatment of chronic inflammatory diseases.

# 35 FEATURES OF PROTEIN ENCODED BY GENE NO: 180

This gene is expressed several infant tissues including brain and liver and various adult tissues including brain, lung, liver, testes, and prostate.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of diseases and conditions which include, but are not limited to, brain cancer, lung cancer, liver cancer and cancers of the reproductive system. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, hepatic system, and reproductive system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., brain and other tissue of the nervous system, lung, liver, testes and other reproductive tissue, and prostate, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an

The tissue distribution of this gene product indicates that the protein product of this clone is involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

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individual not having the disorder.

# 20 FEATURES OF PROTEIN ENCODED BY GENE NO: 181

This gene is expressed primarily in activated monocytes and to a lesser extent in melanocytes and dendritic cells.

Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of immune system diseases and cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the immune system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., blood cells, melanocytes, and dendritic cells, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

35 The tissue distribution indicates that the protein product of this clone could be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 182

This gene is expressed primarily in placenta and several tumors of various tissue origin and to a lesser extent in normal tissues including liver, lung, brain, and skin,

gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an biological sample and for diagnosis of cancers of all kinds. Similarly, polypeptides and issues of the nervous system, and skin, and cancerous and wounded tissues) or bodily antibodies directed to these polypeptides are useful in providing immunological probes system and skin, expression of this gene at significantly higher or lower levels may be for differential identification of the tissue(s) or cell type(s). For a number of disorders fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard routinely detected in certain tissues and cell types (e.g., liver, lung, brain and other of the above tissues or cells, particularly of the central nervous system, respiratory Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a individual not having the disorder.

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activity of the protein product of this clone could be used to block its potential role in The high expression of this gene in multiple tumors indicates that the protein product of the clone may be involved in cell growth control and therefore would be useful for treatment of certain cancers. Likewise molecules developed to block the tumor growth promotion.

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## FEATURES OF PROTEIN ENCODED BY GENE NO: 183

The translation product of this gene shares sequence homology with the mouse This gene is expressed multiple cell types and tissues including brain, lung, Ndr1 gene which is thought to be important in cancer progression. 22

kidney, bone marrow, liver, and spleen.

biological sample and for diagnosis of all types of cancers. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a

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systems, expression of this gene at significantly higher or lower levels may be routinely system, lung, kidney, bone marrow, liver and spleen, and cancerous and wounded detected in certain tissues and cell types (e.g., brain and other tissue of the nervous of the above tissues or cells, particularly of the nervous, immune, and endocrine 35

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another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or fluid from an individual not having the disorder.

molecules developed to block the activity of the protein product of this clone could be The tissue distribution and homology to Ndr1 gene, which is thought to be corresponding to this gene are useful for treatment of certain cancers. Likewise involved in cancer progression, indicate that polynucleotides and polypeptides used to block its potential role in tumor growth promotion. Ś

## FEATURES OF PROTEIN ENCODED BY GENE NO: 184

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This gene is expressed primarily in early stage human brain and liver and to a lesser extent in several other fetal tissues.

biological sample and for diagnosis of diseases and conditions; brain and liver cancers. wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal brain and other tissue of the nervous system, liver, and fetal tissue, and cancerous and fluid) or another tissue or cell sample taken from an individual having such a disorder, higher or lower levels may be routinely detected in certain tissues and cell types (e.g., providing immunological probes for differential identification of the tissue( $ec{s}$ ) or cell central nervous system and immune system, expression of this gene at significantly Similarly, polypeptides and antibodies directed to these polypeptides are useful in lype(s). For a number of disorders of the above tissues or cells, particularly of the Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 2 ន 25

relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

The expression of this gene in embryonic tissues indicates that the protein could be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 185

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This gene is expressed primarily in infant and embryonic brain.

biological sample and for diagnosis of degenerative nervous system disorders and brain cancer. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a 33

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type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., embryonic tissue, brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the disorder.

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The expression of this gene in embryonic tissues indicates that the protein could 10 be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

## FEATURES OF PROTEIN ENCODED BY GENE NO: 186

This gene is expressed primarily in multiple tissues including placenta, fetal lung, fetal liver, and brain.

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Therefore, polynucleotides and polypeptides of the invention are useful as reagents for differential identification of the tissue(s) or cell type(s) present in a biological sample and for diagnosis of all types of cancers including liver, brain and lung. Similarly, polypeptides and antibodies directed to these polypeptides are useful in providing immunological probes for differential identification of the tissue(s) or cell type(s). For a number of disorders of the above tissues or cells, particularly of the central nervous system, pulmonary system, and hepatic system, expression of this gene at significantly higher or lower levels may be routinely detected in certain tissues and cell types (e.g., placenta, lung, liver, and brain and other tissue of the nervous system, and cancerous and wounded tissues) or bodily fluids (e.g., serum, plasma, urine, synovial fluid or spinal fluid) or another tissue or cell sample taken from an individual having such a disorder, relative to the standard gene expression level, i.e., the expression level in healthy tissue or bodily fluid from an individual not having the

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The expression of this gene in embryonic tissues indicates that the protein could be involved in growth regulation and could be used as a growth factor or growth blocker in a variety of settings including treatment of cancers.

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ DO: N	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	Start Codon	AA of Signal Pep	NO: Y	First AA of Sig Pep	AA of Sig Pep		AA of ORF
1	HTTEZ21	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	11	582	1	582	177	177	313	1	18	19	22
1	HTTEZ21	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	197	1020	296	830	442	442	499	1	18	19	22
2	HBGBW52	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	12	465	1	465	81	81	314	1	30	31	128
2	HBGBW52	97897 02/26/97 209043 05/15/97		198	524	229	343		196	500	1	20	21	33
3	HCUFM41	97897 02/26/97 209043 05/15/97	ZAP Express	13	474	1	474	1	1	315		24	25	28
3	HCUFM41	97897 02/26/97 209043 05/15/97	ZAP Express	199	332	1	319	35	35	501	- 1	24	25	28

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N.	961	18	30	Ī	354	<del>1</del> 9	<del>1/</del> 9	£59	1	£\$9	77	Dogod	L6/SI/SO E40607 L6/9Z/ZO L6/9Z/GO	HCFAW04	71
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	<del>1</del> 9	18	30	Ī	373	<b>7</b> 9	<b>7</b> 9	166	Ī	166	17	AX 4A∑-inU	L6/S1/S0 E4060Z L6/9Z/Z0 L68L6	ниспев	II.
118	<u>L</u> Z			I	322	061	061	17L	SS	ΙÞL	07	AX 4AS-inU	26/51/50 26/97/70 26/97/70 26/97/50	HSAXR76	10
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88	izasi AA Io FISIO	Predicted As triff To Secreted Portion		First AA o o' Sig Pep	SEQ ID	30	S' NT of Start Codon	3' NT of Clone Seq.	5' NT of Clone Seq.	Total NT Seq.	£ÿAö×	Vector	ATCC Deposit No: X and Date	cDNA Clone ID	Gene No.
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Gene : No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgegz x	Total NT Seq.	Seg.	of Clone Seq.	5' NT of Start Codon	AA of Signal Pep	SEQ BO: Y	AA of Sig Pep		Predicted First AA of Secreted Portion	
13	HLMAV65	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	23	1486	596	1418	102	102	325	1			
13	HLMAV65	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	203	847	1	839	87	87	505		30	31	75
13	HLMAV65	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	204	852	75	850		690	506	1			10
13	HTXEF04	209235 09/04/97	Uni-ZAP XR	205	1354	54	1354	100	100	507	1	33	34	207
14	HPMFD84	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	24	2323	1017		1242	1242	326		21	22	68
14	HPMFD84	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	206	1378	113	1226	303	303	508		25	26	36
15	HE6DB26	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	25	683	1	683	304	304	327	1	30	31	84

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ZEQ BO: X	Total NT Seq.	Seq.	of Clone Seq.	5' NT of Start Codon	•	SEQ NO: Y		AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
15	HE6DB26	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	207	1166	281	884	567	567	509	1	18	19	19
16	HHFFL33	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	26	2036	14	1959	214	214	328	1	20	21	36
17	HODBD33	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	27	717	1	717	70	70	329	1	30	31	63
17	HODBD33	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	208	697	2	697	33	33	510	1	31	32	32
18	HMDAE90	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	28	495	1	495	39	39	330	1	24	25	35
19	HOUAW01	97897 02/26/97 209043 05/15/97	Uni-ZAP XR	29	556	1	556	116	116	331	1	19	20	23

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87	87	LZ	ī	115	<b>∠8</b> €	785	756	\$LZ	726	607	pSportl	26/51/50 57/50/607 26/97/70 26826		17
111	31	30	ī	555	£8	<b>48</b>	SIL	1	SIL	18	pSport	26/51/50 5 <del>0</del> 607 26/92/70 26826		17
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izeJ AA Io IGF	Predicted AA tzri-T to fo Secreted notion		First AA To Sig Geg	EO SEO	YN 'S To Isriff To AA Signal GeP	S' NT Io Start Codon	of Clone Seq.	S' NT of Clone Seq.	Total NT Seq.	ZEO NO: XEO		ATCC Deposit No: Z and Date	cDNA Clone ID	Gene No.

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	-	ATCC Deposit		NT SEQ ID	Total NT	of	3' NT of Clone Sea.		5' NT of First AA of Signal	SEQ ID		AA of	Predicted First AA of Secreted	
Gene No.	cDNA Clone ID	No: Z and Date	Vector	NO: X	Seq.	sey.		Codon	Pep	Y	Pep	Pep	Portion	ORF
28	HSSDM73	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	38	672	1	672	22	22	340	1	38	39	42
29	HBMVK68	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	39	1908	135	1908		309	341	1	20	21	26
30	HMKDC66	97898 02/26/97 209044 05/15/97	pSport1	40	458	93	458	147	147	342	1	24	25	26
31	HMKCU94	97898 02/26/97 209044 05/15/97	pSport1	41	1153	500	1153	427	427	343	1	30	31	157
31	HMKCU94	97898 02/26/97 209044 05/15/97	pSport1	213	1079	502	896		739	515	1	23	24	43
32	HRDEW41	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	42	1983	1092	1983	27	27	344	1	11	12	520

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ESES:×	Total NT Seq.	5' NT of Clone Seq.	of	5' NT	5' NT of First AA of Signal Pep 2030	AA SEQ ID	First AA of Sig Pep	AA of	Predicted First AA of Secreted Portion	
32	HRDEW41	97898 02/26/97 209044 05/15/97	Uni-ZAP XR		3791							10	20	39
33	HTOJN06	97898 02/26/97 209044 05/15/97	Uni-ZAP XR		1406	1	695		19	345	1	19		
34	HBGDA21	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	44	1391	851	1153	74	74	346	1	30	31	234
34	HBGDA21	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	215	1334	822	1036		638	517	1	18	19	174
35	HFGAK75		Uni-ZAP XR	45	1569	768	1569	14	14	347	1	19	20	169
35	HFGAK75	97898 02/26/97 209044 05/15/97	Uni-ZAP XR	216	1511	770	1404	844	844	518	1	32	33	43

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44	HMABL38	97899	Uni-ZAP XR	220	1258	149	1190	254	254	522	1	18	19 .	26
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		209045 05/15/97						l		l	1	1		
45	HSKDK47	97899	Uni-ZAP XR	55	1896	596	1614	650	650	357	1	33	34	47
43	HSKUK41	02/26/97	Om-2211 7110	"	.0,0	""					i -			
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46	HOSFH03	97899	Uni-ZAP XR	56	1753	555	1753	414	414	358	1	18	19	73
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47	HOGAV75	97899	pCMVSport	57	1220	690	1024	128	128	359	1	30	31	102
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47	HOGAV75	97899 02/26/97	2.0	1 222	1170	1 ′ 12	1,103	1	1057	1	<b>'</b>	1		Ι΄΄
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zeee Eeeee	Total NT Seq.	5' NT of Clone Seq.	of	5' NT of Start Codon		SEQ ID		AA of	of Secreted	Last AA of ORF
48	HFCAI74	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	58	1049	362	1049	335	335	360	1	33	34	48
49	HAGBI17	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	59	1776	854	1737	189	189	361	1	30	31	179
49	HAGB117	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	223	1791	979	1791	1164	1164	525	1	18	19	40
50	HLFBC91	97899 02/26/97 209045 05/15/97	pBluescript SK-	60	443	Î	443	164	164	362	1	21	22	25
51	HPRCA31	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	61	2888	1909	2888	90	90	363	1	30	31	224
51	HPRCA31	97899 02/26/97 209045 05/15/97	Uni-ZAP XR	224	2517	1597	2517	1953	1953	526	1	18	19	57

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Last AA lo TAO

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81

Last AA of Sig Pep

Secreted Portion

AA izriT lo

Predicted

<u>0LE</u>

69£

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625

**≰**8693≻

AA Ao To gi2 gi2

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<del>19</del>7

193

976

Pep

Signal

30 AA Jeni 🗗

jo LN S

1562 530

77

56

<del>797</del>

163

976

Codon

lo nai2

ZN .S

805

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988

679

869

9881

Sed.

3' NT Of Clone

808

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679

Uni-ZAP XR 228 2043 1133 1756 1262

732

9881

Total NT Seq.

Sed.

S' NT of Clone

SLE ISLI

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NA GAS-inU

AS abdmal

RX 4AZ-inU

AX 4AZ-inU

AX 4AZ-inU

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68	52	54	L	228	436	984	1033	747	1080	977	AX 4AS-inU	66876	MADJ02	<b>₽</b> \$
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751	31	30	ī	99€	576	576	£88	LEZ	€88	t9	Jai-ZAP XR		TOTA MH	<b>₽</b> \$
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L97	56	52	١,	392	<del>1</del> 96	<del>1</del> 96	3492	£88	3245	£9	ssənqx3 4AS	66876	HHTLC66	£\$
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ORF	Portion	Ьeр	454	,	Pep	Codon			Seq.	х	Vector	and Date	Clone ID	.oN
ìo	Secreted		BiZ		Signal	Start	Seq.	Sed.	TN	:ÔN		Z:0N	ANGS	SnsD
ΑA	ìo	10	30	OI	30 AA	10	Clone		IstoT	Œ		Deposit		ı
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L6/97/70 668L6

76/51/50 5<del>0</del>607 L6/97/70

66816 668740 76/21/20 76/32/20 76/32/20

76/82/<del>1</del>0 110602

02\12\6\ \$0607 *L6/97/7*0 668*L*6

26/51/50 50607 26/97/70 66826

L6/\$1/\$0 209045 76/92/20

66876

and Date

Deposit No: Z

**DOTA** 

HCMEE<sub>00</sub>

HHFHN61

**HWEEX43** 

**HKTAG33** 

**59SXA2H** 

**ESEXASH** 

**НР**ВСО93

Clone ID

**VND** 

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85

95

95

Gene No.

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgagx	NT Seq.	of Clone Seq.	Seq.	5' NT		SEQ NO: Y	AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
59	HCWEF90	97899 02/26/97 209045 05/15/97	ZAP Express		448	9	448		. 1	532	1	22	23	75
60	HHGCM20	97899 02/26/97 209045 05/15/97	Lambda ZAP II	70	245	1	245	93	93	372	1	1	2	51
61	HFRAU10	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	71	361	1	361	1	1	373	1	30	31	61
61	HFRAU10	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	231	407	1	407	210	210	533	1	17	18	60
62	HATDT67	97900 02/26/97 209046 05/15/97	Uni-ZAP XR		713	8	713	169	169	374	1	30	31	40
62	HATDT67	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	232	830	190	580	329	329	534	1	28	29	39

Gene No.	cDNA Clone ID HOUBG93	ATCC Deposit No: Z and Date	Vector Uni-ZAP XR	75 E E E E E E E E E E E E E E E E E E E	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT	5' NT of First AA of Signal Pep	SEQ ID	First AA of Sig Pep	Last AA of Sig Pep	of Secreted	
		02/26/97 209046 05/15/97						•05		-50.5				
63	HOUBG93	97900 02/26/97 209046 05/15/97		233	932	138	905	287	287	535	1			2
64	HMWEX24	97900 02/26/97 209046 05/15/97	·	74	4602		4525	730	730	376	1	30	31	203
64	HMWEX24	97900 02/26/97 209046 05/15/97	Uni-Zap XR	234	2786	2406	2739	2577	2577	536	1	22	23	36
65	HSGBA84	97900 02/26/97 209046 05/15/97		75	1255	1	1195		112	377	1	28	29	29
66	HTOCD52	97900 02/26/97 209046 05/15/97		76	475	1	475	13	13	378	1	30	31	136

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Secreted Portion

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¥ SEO NO: VEO VEO

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58

1504

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611

5' NT of First AA of Signal Pep

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611

101

S' NT of Start Codon

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LES

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1024

of Clone Seq.

IN .E

-65

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5024 1609 1953

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Seq.

Clone

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789

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Seq.

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X NO: ID SEO

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Uni-ZAP XR 237 1286

Uni-ZAP XR 83

JAX 4AZ-inU

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AAZ abdma

Vector

JAZ AAZ-inU

AX 4AZ-inU

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133	89 <del>1</del>	16	30	1	380	97	
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	ψĬ			ī	<b>LES</b>	97	
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67	- 21	50	上上	185	<u> 1</u> 97	<i>L</i> 97	8911	9£1	1168	6 <i>L</i>	AX 4AS-inU	00676	HETG109	69
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ψĬ			ī	LES		97	854	1	854		AX 4AS-inU	00020 20807 20807 20907 200626	HTOCDS2	99
128.1 AA 10 190.	Predicted AA izriF o Secreted notion	Last AA of Sig Pep		:ON	for the following the following the following the following to follow the following the follow the follow the follow the follow the follow the follow the follow the follow the follow the follow the follow the follow the follow the follow the following th	S' NT of Start Codon	of Clone Seq.		Total NT Seq.	ZEO NO: NO: X	Vector	ATCC Deposit No: X and Date	cDNA Clone ID	Gene No.

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L6/S1/S0 940607 L6/97/70

209046 26/51/50 9<del>1</del>0607

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26/51/50 9+060Z 26/9Z/Z0 00626

20607 9 L6/97/70

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Gene No.

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	5898;x	NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	SEA SEY	First AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
75	нвіавз9	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	238	734	1	734	1	1	540	1	37	38	108
75	HBIAB39	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	239	809	80	794		294	541	1	15	16	106
76	HTXDU73	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	86	1238	36	918	17	17	388	1			1
77	HOEAS24	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	87	1460	9	1458	166	166	389	1	53	54	299
77	HOEAS24	97900 02/26/97 209046 05/15/97		240	2201	841	2080	507	507	542	1	43	44	136
77	HOEAS24	97900 02/26/97 209046 05/15/97	Uni-ZAP XR	241	1661	311	1520	390	390	543	1	35	36	424

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		ATCC		SEQ		of	of	5' NT		SEQ	AA		First AA	
		Deposit		D	Total	Clone		of	AA of	Ē	of	of	of	AA
Gene	cDNA	No: Z		NO:	NT	Seq.	Seq.	Start	Signal	NO:	Sig	Sig	Secreted	of
No.	Clone ID	and Date	Vector	х	Seq.	1	•	Codon	Pep	Y	Pep		Portion	ORF
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78	HTEIY30	97900	Uni-ZAP XR	88	1395	567	1395	639	639	390	1	36	37	49
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1 1	İ	209046 05/15/97											i .	1 1
	HSKNE46	97900	pBluescript	89	1186	352	1186	540	540	391	<del>                                     </del>	49	50	61
79	HSKNE40	02/26/97	photoescript	07	1100	332	1100	340	340	371	١.	~	1 30	"
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79	HSKNE46	97900	pBluescript	242	1146	329	1146	564	564	544	1	21	22	39
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80	HPMFL27	97900	Uni-ZAP XR	90	1821	1203	1614	1503	1503	392	1	30	31	79
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		209046			1		1	l				1	1	1
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81	HMWDN32	97900	Uni-Zap XR	91	862	253	862	359	359	393	1	32	33	36
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		209046		1			1	i			l	1	1	1 1
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82	HPRAX55	97900	Uni-ZAP XR	92	696	349	סאס	78	ا مح	374	'	30	31	180
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										ľ		740602 76/92/20		
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33	7.1	50		949	161	161	116	7.2	1256	544	AAZ sbdmsJ	10616	HCQAV53	58
751	55	25	١,	<b>46٤</b>	907	907	8491	ī	2203	S6	AX 4AZ-inU	20607 940607	H2DEA53	58
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09	77	17	1	968	S8 <i>L</i>	\$8 <i>L</i>	ZLLI	74Z	<b>⊅</b> LL[	<u></u> \$6	NX 4AS-inU	10646	HEZPL777	<del>7</del> 8
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12			1	362	<i>L</i> 61	<i>L</i> 61	65LI	1	9881	£6	AX 4AS-inU	00676	HHFFW36	٤3
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88	33	32	Ī	545	348	348	1230	592	1320	243	AX AAS-inU	00646	<b>LENAX55</b>	78
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NEQ SED NO N	Total NT Seq.	5' NT of Clone Seq.	of Clone	5' NT of Start Codon	5' NT of First AA of Signal Pep	SEQ ID			Predicted First AA of Secreted Portion	Last AA of ORF
91	HTSEL31	97901 02/26/97 209047 05/15/97	pBluescript	101	1394	608	1346	602	602	403	1	23	24	87
92	HAUBL57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	102	794	1	794	518	518	404	1	30	31	92
92	HAUBL57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	248	1766	42	1766	356	356	550	1	30	31	168
92	HAUBL57	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	249	2664	47	1708		147	551	1	18	19	124
93	HODAS59	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	103	1544	898	1531	975	975	405	1			21
94	НЕ6СТ48	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	104	871	106	871	248	248	406	1	34	35	174

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				NT		5' NT			of				Predicted	
		ATCC		SEQ		of	of	5' NT		SEQ			First AA	
		Deposit		Œ	Total	Clone		of	AA of		of	of Sig	of Secreted	AA of
Gene	cDNA	No: Z	37	NO:	NT	Seq.	Seq.	Start Codon	Signal Pep	NU:	Sig Pep	Pep		ORF
No.	Clone ID	and Date	Vector	Х	Seq.				•	_	Гер			
94	HE6CT48	97901	Uni-ZAP XR	250	865	97	865	258	258	552	1	19	20	177
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95	HMDAA61	97901	Uni-ZAP XR	105	404	1	404	16	16	407	1	21	22	64
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95	HMDAA61	97901	Uni-ZAP XR	251	2082	852	2074	829	829	553	1	22	23	1/2
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96	HAQBK61	97901	Uni-ZAP XR	106	1542	300	1342	122	122	400	١.	1 21	32	200
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96	HAQBK61	97901	Uni-ZAP XK	232	1462	مرد ا	1402	l	ودن	1 224	l '	1,2	1 10	[ " ]
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1-0	INCHINDOL	209215		253	834	1	834	82	82	555	1	40	41	251
96	HCUHB01	08/21/97		1233	054	1 '	۱۳۶۹	62	""	در ا	Ι'.	1 70	71	1231
1-07	TYAODETT	97901	Uni-ZAP XR	107	2327	1528	2327	465	465	409	1	30	31	284
97	HAQBF73	02/26/97		1 '''	12321	1,728	2321	703	1 703	"	.*	1	''	1207
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<b>L17</b>	1£	30	I	LSS	9 <i>L</i> I	941	7431	765	5214	552	AX 4AZ-inU	26/51/50 25/5007 26/97/70 10626	<b>НЕТНЕ</b> 07	66
752	ΙE	30	I	111	£06	£06	1052	SLZ	6887		AX 4AZ-inU	L6/S1/S0 L1 <del>0</del> 0607 L6/97/70 106L6	то <u>антан</u>	66
781	58	82	ī	0110	2/1	2/1	7901	<b>LS1</b>	7901	801	AX 4AS-inU	L6/S1/S0 L <del>1</del> 0607 L6/97/70 106L6	<b>46ТВОАН</b>	86
61			ı	955	886		8051	588	80SI			6/51/50 6/97/70 106/6	<b>ЕГ</b> ЧВОАН	<i>L</i> 6
Last AA Io SRF	AA 12117 To Secreted Portion	AA To Sig Goq	AA of Sig qaq		first To AA Signal Pep	S' NT of Start Codon	of Clone Seq.	of Clone Seq.	Total NT Seq.	X NO: ZEO	Vector	ATCC Deposit No: Z and Date	cDNA Clone ID	Gene No.
	Predicted	126.J	First		Jo .		JN .E	ZN .S		TN				ŀ

. 693															
PCT/US98/04493													L6/S1/S0 L <del>1</del> 0607 L6/97/70		
G	_35_			I	-517-	-60 <i>L</i> -	60 <i>L</i>	- <del>1</del> 591-	-655-	-Þ\$91-	-6-1-1-	-NX-AAS-inU-		HEBD185-	£01-
Ã.	· 0s	9£	32	ī	795	SE	SE	1562	97	1562	. 097	AX 4AS-iāU	L6/51/50 L70607 L6/97/70 106L6	HEZBG03	201
	15	16	30	Ī	blb	172	172	EIEI	871	EIEI		AX 4AZ-inU	L6/\$1/\$0 L <del>0</del> 607 L6/97/70	НЕЅВСОЗ	701
	801	57	54	ī	195	747	747	SEII	69	£611	526	Other	05/15/88 209657	нівекіе	101
142	1.7	07			CYA	T		,,,,,	•	<b>/111</b>		AX 4AZ-inU	26/51/50 26/97/70 26/97/70	HCRAM28	101
	51 24	07 67	61 87	I	220	72	72	<u> </u>	ç	LLEZ		Inoq2q	200119 500110	HEONN28	001
	56	61	81	ı	655	31¢	314	559	218	689	LSZ	9AS sbdmsJ II	76/51/50 76/92/20 76/92/20	HLQAB52	001
WO 98/39448	128.I AA 10 10 FIRO	Predicted AA izriH of Secreted Fortion	AA Io gi2 gaq	tzriH AA To giS gsA	<b>S</b> A Si≻	lengi2 Signal qsq	S' NT of Start Codon	of Clone Seq.	Seq.	Total NT Seq.	NT SEQ NO: NO: X	Vector	ATCC Deposit No: S and Date	cDNA Clone ID	Jene No.
WO 98															

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ NO: N	Total NT Seq.	Seq.	of Clone Seq.	Start Codon	AA of Signal Pep	SEQ ID NO: Y	AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
104	HCUBC79	97901 02/26/97 209047 05/15/97	ZAP Express			540	1171	337	337	416	1	30	31	163
104	HCUBC79	97901 02/26/97 209047 05/15/97	ZAP Express	261		626	1161	335	335	563	1	30	31	253
104	HCUBC79	97901 02/26/97 209047 05/15/97	ZAP Express	262	1162	629	1131	942	942	564	1			18
105	HSVAF07	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	115	842	373	800	100	100	417	1	65	66	174
105	HSVAF07	97901 02/26/97 209047 05/15/97	Uni-ZAP XR		735	290	735			565	1			
105	HSVAF07	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	264	783	416	783		413	566	1	33	34	73

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ZEQ SEQ DO: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon		SEQ ID NO: Y			Predicted First AA of Secreted Portion	Last AA of ORF
106	НТЗАМ65	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	116		187	1470	581	581	418	1		31	263
106	НТЗАМ65	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	265	1638	301	1405	119	119	567	1	30		
106	НТЗАМ65	97901 02/26/97 209047 05/15/97		266	1455	148	1188	438	438	568	1	24	25	70
107	HE6DK18	97901 02/26/97 209047 05/15/97			952	418	906	499	499	419		28	29	120
108	НЕВЕК93	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	118	1256		1079		301	420		30	31	159
108	НЕВЕК93	97901 02/26/97 209047 05/15/97	Uni-ZAP XR	267	1086	25	1050	227	227	569	1	23	24	34

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Last AA lo 190

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Secreted Portion

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Last AA of Sig Pep First AA To Sig Pep 715

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-087

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Signal qsq

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143

S' NT of Start Codon

**LE9** 

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1910

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9L61

of Clone Seq.

LN .8

LL

688

6L11

581

Seq.

To SnolD

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1350

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1221

1802

256

Total NT Seq.

124 1717

1/2

X NO: SEO

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146

87	LZ	I	ZLS	LEE	LEE	995	ī	†LS	0/2	AX 4AS-inU	L6/\$1/\$0 L70607 L6/97/70 106L6	HOEVM81	111
18	30	I	£ZÞ	oş	OS.	609	81	019	121		L6/S1/S0 L40607 L6/97/70 106L6	HOEVM81	111
££	35	I	777	881	861	07/1	ī	7871	150	AX 4AZ-inU	26/51/50 20007 26/97/70 10626	нахвглях Нахви	011
87	LT	ī	115	757	757	5101	<b>⊅</b> ∠1	1534	697		26/51/50 25/0607 26/97/70 10626	нъсміо	601
32	34	ī	0/5	SII	SII	1003	17	1003	897	AX 4AZ-inU	26/51/50 26/97/70 26/97/70	HIPCM10	601
ıç	os	ī	174	SLI	SLI	1501	141				L6/97/70 L6/97/70 L6/97/70	HIPCM10	601
AA ızıi7 10	AA	First AA To Sig Pep	ID ZEÓ	to teriT to AA	TN 'S lo hal	of Clone Seq.	of Clone Seq	Total NT Seq.	E S S S S S S S S S S S S S S S S S S S	Vector	ATCC Deposit No: Z and Date	cDNA Clone ID	Gene No.
	First AA of of Secreted Portion 35 31 35 31	30 31 35 33 37 38 34 32 34 32 34 32 34 32	AA AA First AA of of of of of of of of of of of of of	A22 I 30 31  A21 I 20 31  A21 I 20 31  A21 I 20 31  A21 I 34 35  A21 I 30 31  A21 I 30 31	10   10   10   10   10   10   10   10	20 20 423 I 30 31  138 138 422 I 30 31  137 112 210 1 34 35  138 138 423 I 30 31  138 138 423 I 30 31  149 140 15 15 25 25  15 210 210 21 34 35  16 210 210 21 34 35  17 21 21 21 21 21 21 21  18 21 21 21 21 21 21  21 21 21 21 21 21  22 23 21 21 21 21 21  23 21 21 21 21 21 21  24 25 21 21 21 21 21 21 21 21 21 21 21 21 21	1003 115 175	18 609 50 50 423 1 30 31  194 1015 232 232 571 1 27 28  171 1063 115 115 217 21	1782   1   1720   138   138   422   1   30   31   31   32   33   34   34   35   34   35   35   35	120   1782   1   1720   138   138   422   1   30   31   31   32   33   33   34   34   34   34   35   34   35   35	Uni-ZAP XR 120 1782 1 1720 138 138 422 1 30 31  Uni-ZAP XR 120 1782 1 1720 138 138 422 1 30 31  Uni-ZAP XR 120 1782 1 1720 138 138 422 1 32 33  Uni-ZAP XR 120 1782 1 1720 138 138 422 1 32 33  Uni-ZAP XR 269 1234 174 1015 232 232 571 1 27 28  Uni-ZAP XR 169 1143 174 1015 232 232 571 1 27 28  Uni-ZAP XR 269 1234 174 1015 232 232 571 1 20 31  Uni-ZAP XR 269 1234 174 1015 232 232 571 1 30 31  Uni-ZAP XR 269 1234 174 1015 232 232 571 1 30 31  Uni-ZAP XR 269 1234 174 1015 232 232 571 1 30 31  Uni-ZAP XR 269 1234 174 1015 232 232 571 1 30 31	ATICC Deposit No. Z	HOEAWSI 970047   ATCC

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L6/51/50 8+0607 L6/97/70 706L6 L6/51/50 8+0607 L6/97/70 706L6

706L6 L6/S1/S0 L4/9607 L6/97/70 106L6 L6/S1/S0 L4/0607 L6/97/70 L4/0607 L4/0607 L4/0607 L4/0607 L4/0607

ATCC Deposit No: X and Date

HOVBA03

HOVBA03

HTXGS75

HEAAR60

HEAAR60

HOEV<sub>b</sub>¢1

cDNA Clone ID

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SII

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Сепе Ио.

Inoq2q

Inoq2q

AX 4AZ-inU

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Vector

WO 98/39448

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WO 98/39448

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zgag×	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	AA of Signal Pep	SEQ NO: Y		AA of Sig Pep	of Secreted Portion	Last AA of ORF
116	HGBGK76	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	126	431	1	431	73	73	428	1	38	39	47
116	HGBGK76	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		515	1	515	43	43	-575	1	20	21	30
117	HBMUW78	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		3752	3465	3752	748	748	429	1	30	31	370
117	HBMUW78	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		2995	2738	2995	2777	2777		1	18	19	29
118	HASAS24	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		1144	669	1144	896	896	430	1			30
119	HSIDN55	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	129	1830	1234	1830	1265	1265	431	1			24

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ZEESX ZEESX	Total NT Seq.	Seq.	of Clone Seq.	5' NT of Start Codon		SEQ NO: Y	First AA of Sig Pep	AA of Sig Pep	Portion	Last AA of ORF
120	HGBGZ64	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	130	1864	1505	1741	1578	1578	432	1	37	38	53
121	Н6ЕВЈ64	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	131	2041	1	1214	46	46	433	1	35	36	176
121	Н6ЕВЈ64	97902 02/26/97 209048 05/15/97		275	1990	8	1128	71	71	577	1	16	17	92
122	HOECP43	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	132	2012	853	1986	1127	1127	434	1	22	23	77
123	H2CBV31	97902 02/26/97 209048 05/15/97		133	1669		1632		962	435	1	25	26	32
124	HPCAD23	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	134	1565	281	1565	274	274	436	1	25	26	30

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53	97	52	1	144	285	585	LVEI	ZLS	9441	681	AX 4AZ-inU		443СЛ65	671
							27.0.		2111	00,	4214211	L6/\$1/\$0	771	
	l i											209048	-	
			_ 1									<i>L6/97/70</i>		
33	61	81		044	£811	1183	76L1	1044	2561	138	Inoq2q	Z06L6	SEUAY TH	178
1								1				L6/S1/S0		
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77	31	οε	וו	872	97 <i>L</i>	97 <i>L</i>	9642	ZLS	2436	9/2	AZ sbdms.	70676	8⊅HHS∩H	<i>L</i> Z1
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1					1							L6/ST/S0		
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61			τ	857	<b>LOI</b>	<b>LOI</b>	0811	1	1671	136	AX 4AS-inU	70676	негонзт	156
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69	07	6٤	Щ,	LEP	1154	1154	7002	1011	2002	581	Inoq2q	706 <u>L6</u>	RIDAGEH	152
ORF	Portion	Pep	Lep	X.	454	Codon			Seq.	x	Vector	and Date	Clone ID	.oN
10	Secreted	giZ	giZ	ЮN	Lsngi2	Start	Seq.	Seq.	IN	:ON	1	Z:ON	ANGS	Gene
₩	30	ìo	ÌO	ID.	30 AA	30	Clone		LetoT	ID,	1	Deposit		1
126.J	AA ızıiH	AA	ΑA	SEO	121iH	TN 'S	10	10		SEQ	İ	DDTA		
1	benoiberq	Last	triiA	ΑA	30		TN 'E	TN 'S		TN	ļ			
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HPMGD24

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Gene No.

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206.26 206.26 206.26 206.26 206.26 206.26 206.26

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ATCC Deposit No: Z and Date

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Secreted Portion

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of First AA of Signal Pep

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Vector

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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ NO: N	Total NT Seq.	Seq.	of Clone Seq.	Start Codon	of First AA of Signal Pep	SEQ NO: Y	First AA of Sig Pep	AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
135	HPTVC60	97902 02/26/97 209048 05/15/97	pBluescript	145	1021	526	1021	74	74	447	1	30	31	278
135	HPTVC60	97902 02/26/97 209048 05/15/97	pBluescript	278	961	524	961	545	545	580	1	23	24	110
136	HSKNE18	97902 02/26/97 209048 05/15/97	pBluescript	146	1285	5	1285	116	116	448	1	30	31	199
136	HSKNE18	97902 02/26/97 209048 05/15/97	pBluescript	279	1228	9	1228	324	324	581	1	26	27	30
137	HMWIF35	97902 02/26/97 209048 05/15/97	Uni-Zap XR	147	1386	169	1272	165	165	449	1	30	31	258
137	HMWIF35	97902 02/26/97 209048 05/15/97	Uni-Zap XR	280	1327	169	1208	160	160	582	1	23	24	71

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	ZT SEQ DO: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	AA of Signal Pep	SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ	First AA of Sig Pep	AA of Sig Pep	of Secreted Portion	AA of ORF
138	HMWGI25	97902 02/26/97 209048 05/15/97	Uni-Zap XR	148	2098	721	2044	784	784	450	1	18	19	87
139	HSKGF03	97902 02/26/97 209048 05/15/97	pBluescript	149	1847	1689	1847	241	241	451	)	33	34	315
139	HSKGF03	97902 02/26/97 209048 05/15/97	pBluescript	281	799	1	799		243	583	1	12	13	47
140	HMSKE75	97902 02/26/97 209048 05/15/97	Uni-ZAP XR			113	1517	417	417	452	1	21	22	52
141	нсмѕн30	97902 02/26/97 209048 05/15/97	Uni-ZAP XR		1540	538	1540	48	48	453	1	30	31	383
141	HCMSH30	97902 02/26/97 209048 05/15/97	Uni-ZAP XR	282	2196	270	2196	294	294	584		32	33	39

493		· · · · · · · · · · · · · · · · · · ·											<i>L6/</i> 51/50		
PCT/US98/04493												II	6 <del>1</del> 0607 66/97/70		
. 107	23	7.1	70	1	985	878	878	1482	912	1634	787	AAZ spdmsJ	£0676	HEXHL79	571
. LC													<i>L6/\$1/\$</i> 0		
					•							11	6 <del>1</del> 0607 26/97/70		
	501	_ 5¢ _	53	1	LSt	1117	111	1502	699	1602	SSI	AAZ sbdmsJ	£06L6	<b>6</b> ГЛНХЫН	571
													<b>L6/S1/S0</b>		
													209048		
	97	77	77	,	954	012	0 <del>1</del>	215		1011	+CI	AX 4AZ-inU	<i>L6/97/7</i> 0 706 <i>L</i> 6	HFAMGI3	144
					231	_U/	_Ur_	CIS		1011	121	ax avz :-11	<i>L</i> 6/\$1/\$0	HEVIOLIS	- //-
		1											2090 <del>4</del> 8		
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153	61				285	179	179	9911	LLZ	2811	283	pBluescript	706L6	HBMDW46	143
													709048 8 <del>1</del> 0607		
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	163	LZ	97	1	554	\$61	561	٤98	I	£98	ESI	pBluescript	Z06L6	HBMDW46	143
								]			Į.		L6/ST/SO		
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	981	٤S	25	ī	<b>484</b>	9	9	SLSI	069	6141	125	pSport	70676	HLMCB92	745
	ЭЯО	Рогиоп	Pep	Pep	X	Pep	пороЭ			Seq.	х	Vector	and Date	Clone ID	.oN
	10	Secreted	BiZ	Sig	:ÔN	Isngi2	Start	.ps2	Seq.	TN	:ÔN		Z:ON	4NQ5	эпэД
	₩	10	30	lo	Œ	îo AA	10	Clone			Œ		Deposit		
	126.]	AA mii	₩	ΑA	SEÓ		TN 'S	10	10		SEO		DOTA		
9448		Predicted	ton I	JeniH	AA	TN 'S		TN 'F	IN .S		TN				
ð						JULY 13				L		<u></u>			

76/21/20 6+0607 76/92/20 20646

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05/12/97 209049 709049

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**L6/51/50** 509049 L6/97/70 £06L6

and Date

Deposit No: Z

**DOTA** 

**HCQAV96** 

H2KCO76

HS1AP03

HCFBC03

HCFBC03

HZNYKI

H2NYKI.

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Gene No.

П 4AZ abdma1

pBluescript

AX 9AZ-inU

pSport

Inoq2q

Uni-ZAP XR

AX AAS-inU

Vector

129 2395

858 987

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TV TV

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Portion

Secreted betoiberq AA tzriT Io 23

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Pep

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krift AA To Sig gəq

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6081 8571

3' NT of Clone Seq.

5' NT Of Clone Seq.

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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zees zees zees	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	SEQ ID NO: Y	First AA of Sig Pep	AA of Sig	Predicted First AA of Secreted Portion	Last AA of ORF
150	HSHCC16	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		2120		2108	1416	1416		1			14
151	HTLEF62	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		900	482	900	46	46	463	1	30	31	285
151	HTLEF62	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		1517	783	1517	1062	1062	590	1			24
152	HTLAD94	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	162	1003	1	1003	288	288	464	1	30	31	80
152	HTLAD94	97903 02/26/97 209049 05/15/97	Uni-ZAP XR		3865		1195		281	591	1	16	17	38
153	HTSFQ12	97903 02/26/97 209049 05/15/97	pBluescript	163	2196	1607	2180	1611	1611	465	1	30	31	47

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	を SEQ PS: X	Total NT Seq.	5' NT of Clone Seq.	of Clone Seq.	5' NT of Start Codon		SEQ NO: Y		AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
154	HE6FL83	97903 02/26/97 209049 05/15/97		164	1945	271	1840	299	299	466	1	63	64	96
154	HE6FL83	97903 02/26/97 209049 05/15/97		290	1910	279	1818	355	355	592		39	40	69
155	HTXFJ55	97903 02/26/97 209049 05/15/97		165	2933	489	2871	258	258	467	1	30	31	399
155	HTXFJ55	97903 02/26/97 209049 05/15/97	1	291	3276	486	2838		525	593	1	45	46	308
156	НЈРСЈ76	97903 02/26/97 209049 05/15/97			2243	343	2221		341	468				1
157	HLTED27	97903 02/26/97 209049 05/15/97	i	167	1816	1130	1816	284	284	469	1	31	32	273

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35	54	52	ī	٤٧٦	77.L1	77.LI	2100	7 <del>1</del> 91	2100	141	2K- bBluescript	26/51/50 6†0607 26/97/70 20626	HAWBA28	191
54				S65	015	015	1051	438	1051	263	AX 4AS-inU	20903 670607 20903 20915 20915	HCEUB21	190
102	9 <del>t</del>	SÞ	t	7 <i>L</i> \$	1001	1001	6981	86 <i>L</i>	£881		AX 4A.Z-inU	26/\$1/\$0 6 <del>1</del> 0607 26/97/70 20626	HCEFB51	160
234	LZ	56	ī	14	_61	61	918	94	706	691	pBluescript	26/51/50 61/97/70 20626	HNFIP24	651
761	61	81	I	04	802	802	L8L	Į	S#6	891	Inoq2q	26/\$1/\$0 670607 26/97/70 20626	HWKB¥¢¢	851
77			ı	<b>76</b> 5	90£1	1306	8 <del>1</del> 21	8601	\$691	767	AX 4AS-inU	76/51/50 670607 76/97/20 20676	нстергу	LS1
Last AA Io IS	Predicted AA 12117 To lo Secreted Inotion	AA 10	라 라 아 양양	<b>≰</b> ⋈аў≻	of Sirst To AA Signal Signal Pep	S' NT of Start Codon	ìo	5' NT of Clone Seq.	Total NT Seq.	zgeż×	volceV	ATCC Deposit No: X and Date	cDNA Clone ID	Gene No.
AA 3o	AA 121iT To batataa2	AA 10 3i2	AA Jo Bi2	NO: SEO	121i7 To AA Isngi2	10 Tast 2	of Clone	10 Clone	NT. Seq.	Я В В В В В В В В		Deposit No: N and Date		

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312	15	30	L	SLT	122	155	1571	796	6051	ELI	pBluescript	£0616	ELTATAH	163
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54			,	965	164	1,52	6007				2K-	<i>L6/97/</i> 70		
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ot	Secreted	giZ	Sig	ONI	Signal	Start	Sed.	Seq.	Seq.	X	Vector	and Date	Clone ID	No.
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Last	AA ızıi7	₩	W	2EO	isnii	TN 'S	10	10	10407	SEQ		DOTA		
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	NT SEQ M NO: X		5' NT of Clone Seq.			AA of Signal	SEQ ID		AA of	Predicted First AA of Secreted Portion	Last AA
165	HEAAL31	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	.175	991	374	970	60	60	477	1	24	25	178
165	HEAAL31	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	297	2416	1387	2413	1473	1473	599	1	18	19	25
166	HFKFX55	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	176	1290	499	1290		688	478	1	25	26	52
167	H2LAOI1	97903 02/26/97 209049 05/15/97	pBluescript SK-	177	2290	1	2290	173	173	479	1	22	23	62
168	HPFDZ95	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	178	549	1	549	11	11	480	1	21	22	27
168	HPFDZ95	97903 02/26/97 209049 05/15/97	Uni-ZAP XR	298	545	1	545	17	17	600	1	21	22	27

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	zeg es zeg es ze x	Total NT Seq.	of	3' NT of Clone Seg.	5' NT of Start Codon	5' NT of First AA of Signal Pep	SEQ ID		AA of Sig	Predicted First AA of Secreted Portion	
169	HPTTUII	97904 02/26/97 209050 05/15/97	Uni-ZAP XR		1509	294	1352	92	92	481	1	30	31	339
169	НРТТÜП	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	299	1530	385	1530	562	562	601	1	23	24	61
170	HCFAE79	97904 02/26/97 209050 05/15/97	pSport1	180	1316	985	1250	995	995	482	1	26	27	32
171	HTEDJ34	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	181	777	1	777	51	51	483	1	30	31	48
171	HTEDJ34	97904 02/26/97 209050 05/15/97	Uni-ZAP XR		997	244	997	300	300	602	1	23	24	29
172	HODCW06	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	182	791	1	791	14	14	484		29	30	38

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002	15	30	1	884	LSZ	LSZ	0811	794	2121	981	AX 4AS-inU	26/51/50 050607 26/97/70	]	941
54	10	6	1	109	09		9461	7	6957	302		L6/\$1/\$0 0\$0607 L6/97/70 \$06L6		S41
752	15	30	ı	<u> </u>	<u> </u>	<i>L</i> 9	8877	SSE	2562	281	XX 4AZ-inU	L6/S1/S0 0S0607 L6/97/70		SLI
69	LS	95	ı	E09	233	233	5787	SL	2345		ZK- bpinescubt	76/51/50 050602 76/92/20	H2MBF44	<i>\$2.</i>
346	52	. 77	ı	987	181	121	9651	SL	9651	184	2K- bgjnescubt	26/51/50 050607 26/97/70	H2MBF44	<b>⊅</b> ∠1
19	21	07	ı	584	SLS	S <i>L</i> S	1402	946		£81	AX 4AΣ-iπU	26/51/50 250607 26/97/70	HFTAR26	εLì
1285.] AA 10 19(0	Predicted AA izriF To Societed Societed Portion	AA ³o	AA 3i2	:ON	70 'C of first AA of Signal Gep GeP	TN '2	ìo	S' NT Of Clone Seq.	Total NT Seq.	EØAö×	Vector	ATCC Deposit No: Z and Date	cDNA Clone ID	Gene No.

	_57_				-161-	-107-		-100-	_1.07_		-601-		26/51/50 26/97/70 26/97/70		
	-30-		-		-100-	-IUV-	_	-189-	-787-	189	081	AX 4AS-inU	+06L6	_YSAQAH_	-6L1
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	23	61	81		<b>L09</b>	₽\$	<i>\$\$</i>	1521	I	£671	302	AX 4AZ-inU	10646	ISVATAH	841
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Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	75 EQ 15 EQ 15 EQ 15 EX	Total NT Seq.	of Clone Seq.	3' NT of Clone Seq.	Start Codon		SEQ PO P		AA of Sig Pep	Predicted First AA of Secreted Portion	Last AA of ORF
180	HCEEK08	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	190	1014	703	1014	360	360	492	1	30	31	159
180	HCEEK08	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	306	577	1	577		175	608	1			6
181	HAFAUI8	97904 02/26/97 209050 05/15/97	pBluescript SK-	191	2779	2207	2630	1153	1153	493	1	30	31	279
181	HAFAU18	97904 02/26/97 209050 05/15/97	pBluescript SK-	307	2860	163	2860	21	21	609	1	30	31	232
181	HAFAUI8	97904 02/26/97 209050 05/15/97	pBluescript SK-	308	876	275	876	302	302	610	1	32	33	34
182	НЕТВУ74	97904 02/26/97 209050 05/15/97	Uni-ZAP XR	192	1923	30	1923	45	45	494	1	33	34	193

Gene No.	cDNA Clone ID	ATCC Deposit No: Z and Date	Vector	F SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ SEQ	Total NT Seq.	5' NT of Clone Seq.	3' NT of Clone Seq.	5' NT of Start Codon	5' NT of First AA of Signal Pep	SEQ ID	First AA of Sig Pep	AA of Sig	Predicted First AA of Secreted Portion	
183	HTOAF35	97904 02/26/97 209050 05/15/97	Uni-ZAP XR				2286		178	495	1	30	31	205
183	HTOAF35	97904 02/26/97 209050 05/15/97		309	2025	840	2025	971	971	611	1	18	19	21
184	HCRBX32	97904 02/26/97 209050 05/15/97		194	3054	2004	3054		434	496	1	11	12	147
184	HCRBX32	97904 02/26/97 209050 05/15/97		310	3026		3026		2131	612	1			9
185	HEBGB80	97904 02/26/97 209050 05/15/97		195	907	152	907	297	297	497	1	30	31	64
185	HEBGB80	97904 02/26/97 209050 05/15/97		311	712	67	712	107	107	613		18	19	29

12&1 AA of 18O 82

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Last AA of Sig Pep

tzniH AA 10 giZ gaf 86t

NO: NO: SEO VY 572

5' NT of First AA of Signal Pep

12

Secreted Portion

beloiber9 AA lzrii7 lo LZ6

572

S' NT of Start Codon 687

608

3' NT of Clone Seq. S8L

**†**8

of Clone Seq.

LN .S

790

Total NT Seq.

X ID SEO

Vector

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Table I summarizes the information corresponding to each "Gene No." described above. The nucleotide sequence identified as "NT SEQ ID NO.X" was assembled from partially homologous ("overlapping") sequences obtained from the "cDNA clone ID" identified in Table I and, in some cases, from additional related DNA clones. The overlapping sequences were assembled into a single contiguous sequence of high redundancy (usually three to five overlapping sequences at each nucleotide position), resulting in a final sequence identified as SEQ ID NO:X.

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The cDNA Clone ID was deposited on the date and given the corresponding deposit number listed in "ATCC Deposit No.Z and Date." Some of the deposits contain 10 multiple different clones corresponding to the same gene. "Vector" refers to the type of vector contained in the cDNA Clone ID.

"Total NT Seq." refers to the total number of nucleotides in the contig identified by "Gene No." The deposited clone may contain all or most of these sequences, reflected by the nucleotide position indicated as "5" NT of Clone Seq." and the "3" NT of Clone Seq." of SEQ ID NO:X. The nucleotide position of SEQ ID NO:X of the putative start codon (methionine) is identified as "5" NT of Start Codon." Similarly, the nucleotide position of SEQ ID NO:X of the predicted signal sequence is identified as "5" NT of First AA of Signal Pep."

The translated amino acid sequence, beginning with the methionine, is identified
as "AA SEQ ID NO:Y," although other reading frames can also be easily translated
using known molecular biology techniques. The polypeptides produced by these
alternative open reading frames are specifically contemplated by the presentinivention.

The first and last amino acid position of SEQ ID NO:Y of the predicted signal peptide is identified as "First AA of Sig Pep" and "Last AA of Sig Pep." The predicted first amino acid position of SEQ ID NO:Y of the secreted portion is identified as "Predicted First AA of Secreted Portion." Finally, the amino acid position of SEQ ID NO:Y of the last amino acid in the open reading frame is identified as "Last AA of ORF."

Deposit No: Z and Date

**DOTA** 

HFAMH74

Clone ID

981

981

Gene No.

SEQ ID NO:X and the translated SEQ ID NO:Y are sufficiently accurate and

30 otherwise suitable for a variety of uses well known in the art and described further below. For instance, SEQ ID NO:X is useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO:X or the cDNA contained in the deposited clone. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO:Y may be used to generate antibodies which bind specifically to the secreted proteins encoded by the cDNA clones identified in Table 1.

sequence. In these cases, the predicted amino acid sequence diverges from the actual 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion amino acid sequence, even though the generated DNA sequence may be greater than sequencing errors. The errors exist as misidentified nucleotides, or as insertions or in an open reading frame of over 1000 bases) deleted nucleotides cause frame shifts in the reading frames of the predicted amino acid deletions of nucleotides in the generated DNA sequence. The erroneously inserted or Nevertheless, DNA sequences generated by sequencing reactions can contain

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5 8 15 sequence or the amino acid sequence, the present invention provides not only the sequencing the deposited clone in accordance with known methods. The predicted containing a human cDNA of the invention deposited with the ATCC, as set forth in amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA generated nucleotide sequence identified as SEQ ID NO:X and the predicted translated containing the deposited human cDNA, collecting the protein, and determining its determined by peptide sequencing or by expressing the protein in a suitable host cell acid sequence of the protein encoded by a particular clone can also be directly amino acid sequence can then be verified from such deposits. Moreover, the amino Table 1. The nucleotide sequence of each deposited clone can readily be determined by Accordingly, for those applications requiring precision in the nucleotide

identifying or amplifying the corresponding gene from appropriate sources of genomic Such methods include preparing probes or primers from the disclosed sequence and accordance with known methods using the sequence information disclosed herein. SEQ ID NO: Y, or the deposited clone. The corresponding gene can be isolated in The present invention also relates to the genes corresponding to SEQ ID NO:X.

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sequences provided herein and screening a suitable nucleic acid source for the desired homologs may be isolated and identified by making suitable probes or primers from the Also provided in the present invention are species homologs. Species

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polypeptides, synthetically produced polypeptides, or polypeptides produced by a polypeptides include isolated naturally occurring polypeptides, recombinantly produced combination of these methods. Means for preparing such polypeptides are well The polypeptides of the invention can be prepared in any suitable manner. Such

mature form, or may be a part of a larger protein, such as a fusion protein (see below). The polypeptides may be in the form of the secreted protein, including the ઝ

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such as multiple histidine residues, or an additional sequence for stability during secretory or leader sequences, pro-sequences, sequences which aid in purification, recombinant production. It is often advantageous to include an additional amino acid sequence which contains

5 using antibodies of the invention raised against the secreted protein in methods which one-step method described in Smith and Johnson, Gene 67:31-40 (1988) polypeptide, including the secreted polypeptide, can be substantially purified by the form, and preferably are substantially purified. A recombinantly produced version of a are well known in the art. Polypeptides of the invention also can be purified from natural or recombinant sources The polypeptides of the present invention are preferably provided in an isolated

#### Signal Sequences

20 5 cleavage point for that sequence, are available. For instance, the method of McGeoch produce the same predicted cleavage point(s) for a given protein. the range of 75-80%. (von Heinje, supra.) However, the two methods do not always cleavage points of known mammalian secretory proteins for each of these methods is in indicates the amino terminus of the secreted protein. The accuracy of predicting the from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 method of von Heinje, Nucleic Acids Res. 14:4683-4690 (1986) uses the information region and a subsequent uncharged region of the complete (uncleaved) protein. The Virus Res. 3:271-286 (1985), uses the information from a short N-terminal charged Methods for predicting whether a protein has a signal sequence, as well as the

30 shown in Table 1. methods of McGeoch and von Heinje are incorporated. The analysis of the amino acid Engineering 10:1-6 (1997)), which predicts the cellular location of a protein based on was analyzed by a computer program called SignalP (Henrik Nielsen et al., Protein sequences of the secreted proteins described herein by this program provided the results the amino acid sequence. As part of this computational prediction of localization, the In the present case, the deduced amino acid sequence of the secreted polypeptide

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or - 5 residues) of the predicted cleavage point. Similarly, it is also recognized that in shown in SEQ ID NO:Y which have an N-terminus beginning within 5 residues (i.e., + some cases, cleavage of the signal sequence from a secreted protein is not entirely Accordingly, the present invention provides secreted polypeptides having a sequence vary from organism to organism and cannot be predicted with absolute certainty. As one of ordinary skill would appreciate, however, cleavage sites sometimes

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uniform, resulting in more than one secreted species. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

Moreover, the signal sequence identified by the above analysis may not necessarily predict the naturally occurring signal sequence. For example, the naturally occurring signal sequence may be further upstream from the predicted signal sequence. However, it is likely that the predicted signal sequence will be capable of directing the secreted protein to the ER. These polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

## Polynucleotide and Polypeptide Variants

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"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

Applied Math 48:1073 (1988).) Methods commonly employed to determine identity or methods to measure identity between two polynucleotide or polypeptide sequences, the term "identity" is well known to skilled artisans. (Carillo, H., and Lipton, D., SIAM J Press, (1987); and SEQUENCE ANALYSIS PRIMER, Gribskov, M. and Devereux, published techniques. (See, e.g.: (COMPUTATIONAL MOLECULAR BIOLOGY, SEQUENCE ANALYSIS IN MOLECULAR BIOLOGY, von Heinje, G., Academic Research (1984) 12(1):387 (1984)), BLASTP, BLASTN, FASTA (Atschul, S.F. et programs, including the GCG program package (Devereux, J., et al., Nucleic Acids Lesk, A.M., ed., Oxford University Press, New York, (1988); BIOCOMPUTING: INFORMATICS AND GENOME PROJECTS, Smith, D.W., ed., Academic Press, "Identity" per se has an art-recognized meaning and can be calculated using similarity between two sequences include, but are not limited to, those disclosed in "Guide to Huge Computers," Martin J. Bishop, ed., Academic Press, San Diego, New York, (1993); COMPUTER ANALYSIS OF SEQUENCE DATA, PART I, I., eds., M Stockton Press, New York, (1991).) While there exists a number of (1994), and Carillo, H., and Lipton, D., SIAM J Applied Math 48:1073 (1988). Methods for aligning polynucleotides or polypeptides are codified in computer Griffin, A.M., and Griffin, H.G., eds., Humana Press, New Jersey, (1994); 2 20 22 8

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When using any of the sequence alignment programs to determine whether a particular sequence is, for instance, 95% identical to a reference sequence, the parameters are set so that the percentage of identity is calculated over the full length of the reference polynucleotide and that gaps in identity of up to 5% of the total number of nucleotides in the reference polynucleotide are allowed.

sequences or both amino acid sequences. The result of said global sequence alignment Penalty=1, Joining Penalty=30, Randomization Group Length=0, and Cutoff Score=1, sequence (a sequence of the present invention) and a subject sequence, also referred to calculate percent identity and similarity of an amino acid alignment are: Matrix=PAM length in nucleotide bases, whichever is shorter. Preferred parameters employed to (1990).) The term "sequence" includes nucleotide and amino acid sequences. In a Gap Penalty=5, Gap Size Penalty 0.05, and Window Size=500 or query sequence program based on the algorithm of Brutlag et al. (Comp. App. Biosci. 6:237-245 Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty=0.05, and Window 150, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group as a global sequence alignment, can be determined using the FASTDB computer A preferred method for determing the best overall match between a query is in percent identity. Preferred parameters used in a FASTDB search of a DNA sequence to calculate percent identiy are: Matrix=Unitary, k-tuple=4, Mismatch sequence alignment the query and subject sequences are either both nucleotide 2 2 2 As an illustration, a polynucleotide having a nucleotide sequence of at least 95% "identity" to a sequence contained in SEQ ID NO:X or the cDNA contained in the deposited clone, means that the polynucleotide is identical to a sequence contained in SEQ ID NO:X or the cDNA except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the total length (not just within a given 100 nucleotide stretch). In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to SEQ ID NO:X or the deposited clone, up to 5% of the nucleotides in the sequence contained in SEQ ID NO:X or the cDNA can be deleted, inserted, or substituted with other nucleotides. These changes may occur anywhere throughout the polynucleotide.

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Size=500 or query sequence length in amino acid residues, whichever is shorter

Further embodiments of the present invention include polynucleotides having at least 85% identity, more preferably at least 90% identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to a sequence contained in SEQ ID NO:X or the cDNA contained in the deposited clone. Of course, due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the polynucleotides having at least 85%, 90%, 95%, 96%, 97%, 98%, or 99% identity

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575 Science Drive, Madison, WI 53711 (using the local homology algorithm of Smith

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and Waterman, Advances in Applied Mathematics 2:482-489 (1981).)

Package, Version 8 for Unix, Genetics Computer Group, University Research Park,

al., J. Molec. Biol. 215:403 (1990), Bestfit program (Wisconsin Sequence Analysis

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NO:Y or the expressed protein produced by the deposited clone will encode a polypeptide identical to an amino acid sequence contained in SEQ ID

sequence of the polypeptide is identical to the reference polypeptide except that the deleted or substituted with another amino acid, or a number of amino acids up to 5% of polypeptide sequence may include up to five amino acid alterations per each 100 amino reference sequence or in one or more contiguous groups within the reference sequence carboxy terminal positions of the reference amino acid sequence or anywhere between sequence. These alterations of the reference sequence may occur at the amino or the total amino acid residues in the reference sequence may be inserted into the reference acid sequence, up to 5% of the amino acid residues in the reference sequence may be polypeptide having an amino acid sequence at least 95% identical to a reference amino acids of the total length of the reference polypeptide. In other words, to obtain a example, 95% "identity" to a reference polypeptide, is intended that the amino acid those terminal positions, interspersed either individually among residues in the Similarly, by a polypeptide having an amino acid sequence having at least, for

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least 80% identity, more preferably at least 85% identity, more preferably at least 90% biological activity of the protein. the deposited clone. Preferably, the above polypeptides should exhibit at least one amino acid sequence contained in SEQ ID NO: Y or the expressed protein produced by identity, and most preferably at least 95%, 96%, 97%, 98% or 99% identity to an Further embodiments of the present invention include polypeptides having at

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still more preferably at least 96%, 97%, 98%, or 99% similarity to an amino acid deposited clone. sequence contained in SEQ ID NO:Y or the expressed protein produced by the polypeptides having at least 90% similarity, more preferably at least 95% similarity, and In a preferred embodiment, polypeptides of the present invention include

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or both. Especially preferred are polynucleotide variants containing alterations which substitutions due to the degeneracy of the genetic code are preferred. Moreover, produce silent substitutions, additions, or deletions, but do not alter the properties or the human mRNA to those preferred by a bacterial host such as E. coli). combination are also preferred. Polynucleotide variants can be produced for a variety activities of the encoded polypeptide. Nucleotide variants produced by silent of reasons, e.g., to optimize codon expression for a particular host (change codons in variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or added in any The variants may contain alterations in the coding regions, non-coding regions

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several alternate forms of a gene occupying a given locus on a chromosome of an Naturally occurring variants are called "allelic variants," and refer to one of

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techniques or by direct synthesis. Alternatively, non-naturally occurring variants may be produced by mutagenesis allelic variants can vary at either the polynucleotide and/or polypeptide level organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).)

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5 carboxy terminus of this protein. (Dobeli et al., J. Biotechnology 7:199-216 (1988).) exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma (1993), reported variant KGF proteins having heparin binding activity even after loss of biological function. The authors of Ron et al., J. Biol. Chem. 268: 2984-2988 deleted from the N-terminus or C-terminus of the secreted protein without substantial polypeptides of the present invention. For instance, one or more amino acids can be technology, variants may be generated to improve or alter the characteristics of the Using known methods of protein engineering and recombinant DNA

8 2 amino acid position. The investigators found that "[m]ost of the molecule could be the entire length of the molecule. Multiple mutations were examined at every possible 3,500 individual IL-la mutants that averaged 2.5 amino acid changes per variant over analysis of human cytokine IL-1a. They used random mutagenesis to generate over coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational activity similar to that of the naturally occurring protein. For example, Gayle and sequences examined, produced a protein that significantly differed in activity from wild altered with little effect on either [binding or biological activity]." (See, Abstract.) In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide Moreover, ample evidence demonstrates that variants often retain a biological

30 25 are removed from the N-terminus or C-terminus. Whether a particular polypeptide C-terminus of a polypeptide results in modification or loss of one or more biological will likely be retained when less than the majority of the residues of the secreted form deletion variant to induce and/or to bind antibodies which recognize the secreted form functions, other biological activities may still be retained. For example, the ability of a readily be determined by routine methods described herein and otherwise known in the lacking N- or C-terminal residues of a protein retains such immunogenic activities can Furthermore, even if deleting one or more amino acids from the N-terminus or

have little effect on activity. For example, guidance concerning how to make repeats, and substitutions selected according to general rules known in the art so as substantial biological activity. Such variants include deletions, insertions, inversions, Thus, the invention further includes polypeptide variants which show

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phenotypically silent amino acid substitutions is provided in Bowie, J. U. et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies for studying the tolerance of an amino acid sequence to change.

The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

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The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham and Wells, Science 244:1081-1085 (1989).) The resulting mutant molecules can then be tested for biological activity.

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As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asp and Glu; replacement of the samide residues Asp and Glu; Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

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Besides conservative amino acid substitution, variants of the present invention include (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as an IgG Fc fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

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For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).)

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## Polynucleotide and Polypeptide Fragments

In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence contained in the deposited clone or shown in SEQ ID NO:X. The short nucleotide fragments are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt, and even more preferably, at least about 40 nt in length. A fragment "at least 20 nt in length," for example, is intended to include 20 or more contiguous bases from the cDNA sequence contained in the deposited clone or the nucleotide sequence shown in SEQ ID NO:X. These nucleotide fragments are useful as diagnostic probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150, 500, 600, 2000 nucleotides) are preferred.

invention, include, for example, fragments having a sequence from about nucleotide invention, include, for example, fragments having a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250, 251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, and 701 to the end of SEQ ID NO;X or the cDNA contained in the deposited clone. In this context "about" includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini. Preferably, these fragments encode a polypeptide which has biological activity.

In the present invention, a "polypeptide fragment" refers to a short arraino acid sequence contained in SEQ ID NO:Y or encoded by the cDNA contained in the deposited clone. Protein fragments may be "free-standing," or comprised within a larger polypeptide of which the fragment forms a part or region, most preferably as a single continuous region. Representative examples of polypeptide fragments of the invention, include, for example, fragments from about amino acid number 1-20, 21-40, 41-60, 61-80, 81-100, 102-120, 121-140, 141-160, and 161 to the end of the coding region. Moreover, polypeptide fragments can be about 20, 30, 40, 50, 60, 70, 80, 90,

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100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about"

includes the particularly recited ranges, larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes.

Preferred polypeptide fragments include the secreted protein as well as the mature form. Further preferred polypeptide fragments include the secreted protein or the mature form having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of either the secreted polypeptide or the mature form. Similarly, any number of amino acids, ranging from 1-30, can be deleted from the carboxy terminus of the secreted protein or mature form. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotide fragments encoding these polypeptide fragments are also preferred.

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Also preferred are polypeptide and polynucleotide fragments characterized by structural or functional domains, such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions. Polypeptide fragments of SEQ ID NO:Y falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotide fragments encoding these domains are also contemplated.

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Other preferred fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity.

### Epitopes & Antibodies

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In the present invention, "epitopes" refer to polypeptide fragments having antigenic or immunogenic activity in an animal, especially in a human. A preferred embodiment of the present invention relates to a polypeptide fragment comprising an epitope, as well as the polynucleotide encoding this fragment. A region of a protein molecule to which an antibody can bind is defined as an "antigenic epitope." In contrast, an "immunogenic epitope" is defined as a part of a protein that elicits an antibody response. (See, for instance, Geysen et al., Proc. Natl. Acad. Sci. USA

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81:3998-4002 (1983).)

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Fragments which function as epitopes may be produced by any conventional means. (See, e.g., Houghten, R. A., Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985) further described in U.S. Patent No. 4,631,211.)

In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., Cell 37:767-778 (1984); Sutcliffe, J. G. et al., Science 219:660-666 (1983).)

Similarly, immunogenic epitopes can be used to induce antibodies according to methods well known in the art. (See, for instance, Sutcliffe et al., supra; Wilson et al., supra; Chow, M. et al., Proc. Natl. Acad. Sci. USA 82:910-914; and Bittle, F. J. et al., J. Gen. Virol. 66:2347-2354 (1985).) A preferred immunogenic epitope includes the secreted protein. The immunogenic epitopes may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier. However,

sufficient to raise antibodies capable of binding to, at the very least, linear epitopes in a denatured polypeptide (e.g., in Western blotting.)

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is

immunogenic epitopes comprising as few as 8 to 10 amino acids have been shown to be

meant to include intact molecules as well as antibody fragments (such as, for example, Fab and F(ab')2 fragments) which are capable of specifically binding to protein. Fab and F(ab')2 fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation, and may have less non-specific tissue binding than an intact antibody. (Wahl et al., J. Nucl. Med. 24:316-325 (1983).) Thus, these fragments are preferred as well as the products of a FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

#### **Fusion Proteins**

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Any polypeptide of the present invention can be used to generate fusion proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because secreted proteins target cellular locations based on trafficking signals, the polypeptides of the present invention can be

35 locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

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Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur through linker sequences.

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polypeptide to improve stability and persistence during purification from the host cell or preparation of the polypeptide. The addition of peptide moieties to facilitate handling of Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino polypeptide to facilitate purification. Such regions may be removed prior to final acids, particularly charged amino acids, may be added to the N-terminus of the subsequent handling and storage. Also, peptide moieties may be added to the polypeptides are familiar and routine techniques in the art.

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polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) facilitate purification and show an increased half-life in vivo. One reported example immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins Moreover, polypeptides of the present invention, including fragments, and describes chimeric proteins consisting of the first two domains of the human CD4can also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., J. specifically epitopes, can be combined with parts of the constant domain of Biochem. 270:3958-3964 (1995).)

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would be desired. For example, the Fc portion may hinder therapy and diagnosis if the purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D. proteins comprising various portions of constant region of immunoglobulin molecules deleting the Fc part after the fusion protein has been expressed, detected, and purified, Similarly, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion together with another human protein or part thereof. In many cases, the Fc part in a Bennett et al., J. Molecular Recognition 8:52-58 (1995); K. Johanson et al., J. Biol. example, human proteins, such as hIL-5, have been fused with Fc portions for the example, improved pharmacokinetic properties. (EP-A 0232 262.) Alternatively, fusion protein is beneficial in therapy and diagnosis, and thus can result in, for fusion protein is used as an antigen for immunizations. In drug discovery, for Chem. 270:9459-9471 (1995).) 23

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sequences, such as a peptide which facilitates purification of the fused polypeptide. In Moreover, the polypeptides of the present invention can be fused to marker

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derived from the influenza hemagglutinin protein. (Wilson et al., Cell 37:767 (1984).) Chatsworth, CA, 91311), among others, many of which are commercially available. preferred embodiments, the marker amino acid sequence is a hexa-histidine pepude, Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope As described in Gentz et al., Proc. Natl. Acad. Sci. USA 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion protein. such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue,

Thus, any of these above fusions can be engineered using the polynucieotides or the polypeptides of the claimed invention.

## Vectors, Host Cells, and Protein Production

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The present invention also relates to vectors containing the polynucleotide of the vector. Retroviral vectors may be replication competent or replication defective. In the echniques. The vector may be, for example, a phage, plasmid, viral, or retroviral present invention, host cells, and the production of polypeptides by recombinant

as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate, such a virus, it may be packaged in vitro using an appropriate packaging cell line and then

ransduced into host cells.

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latter case, viral propagation generally will occur only in complementing host cells.

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expression constructs will further contain sites for transcription initiation, termination, translation initiating codon at the beginning and a termination codon (UAA, UGA or promoter, such as the phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to and, in the transcribed region, a ribosome binding site for translation. The coding name a few. Other suitable promoters will be known to the skilled artisan. The The polynucleotide insert should be operatively linked to an appropriate portion of the transcripts expressed by the constructs will preferably include a UAG) appropriately positioned at the end of the polypeptide to be translated.

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resistance for eukaryotic cell culture and tetracycline, kanamycin or ampicillin resistance Streptomyces and Salmonella typhimurium cells; fungal cells, such as yeast cells; insect cells such as Drosophila S2 and Spodoptera Sf9 cells; animal cells such as CHO, COS, selectable marker. Such markers include dihydrofolate reductase, G418 or neomycin As indicated, the expression vectors will preferably include at least one appropriate hosts include, but are not limited to, bacterial cells, such as E. coli, genes for culturing in E. coli and other bacteria. Representative examples of

conditions for the above-described host cells are known in the art. 293, and Bowes melanoma cells; and plant cells. Appropriate culture mediums and

pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1 available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, from Pharmacia. Other suitable vectors will be readily apparent to the skilled artisan Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-9

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present invention may in fact be expressed by a host cell lacking a recombinant vector. phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated In Molecular Biology (1986). It is specifically contemplated that the polypeptides of the are described in many standard laboratory manuals, such as Davis et al., Basic Methods transfection, electroporation, transduction, infection, or other methods. Such methods Introduction of the construct into the host cell can be effected by calcium

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chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for phosphocellulose chromatography, hydrophobic interaction chromatography, affinity cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, A polypeptide of this invention can be recovered and purified from recombinant

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tissues and cells, whether directly isolated or cultured; products of chemical synthetic proteins also is efficiently removed in most prokaryotes, for some proteins, this non-glycosylated. In addition, polypeptides of the invention may also include an initial eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and procedures; and products produced by recombinant techniques from a prokaryotic or be recovered from: products purified from natural sources, including bodily fluids, prokaryotic removal process is inefficient, depending on the nature of the amino acid to after translation in all eukaryotic cells. While the N-terminal methionine on most translation initiation codon generally is removed with high efficiency from any protein modified methionine residue, in some cases as a result of host-mediated processes. procedure, the polypeptides of the present invention may be glycosylated or may be mammalian cells. Depending upon the host employed in a recombinant production which the N-terminal methionine is covalently linked Thus, it is well known in the art that the N-terminal methionine encoded by the Polypeptides of the present invention, and preferably the secreted form, can also

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### Uses of the Polynucleotides

reagents. The following description should be considered exemplary and utilizes Each of the polynucleotides identified herein can be used in numerous ways as

S known techniques.

can be used as a chromosome marker polymorphisms), are presently available. Each polynucleotide of the present invention since few chromosome marking reagents, based on actual sequence data (repeat identification. There exists an ongoing need to identify new chromosome markers, The polynucleotides of the present invention are useful for chromosome

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cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the SEQ ID NO:X will yield an amplified fragment. exon in the genomic DNA. These primers are then used for PCR screening of somatic selected using computer analysis so that primers do not span more than one predicted (preferably 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can be Briefly, sequences can be mapped to chromosomes by preparing PCR primers

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be achieved with panels of specific chromosome fragments. Other gene mapping specific-cDNA libraries. sorted chromosomes, and preselection by hybridization to construct chromosome strategies that can be used include in situ hybridization, prescreening with labeled flowpolynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can Similarly, somatic hybrids provide a rapid method of PCR mapping the

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technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This 2,000-4,000 bp are preferred. For a review of this technique, see Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York Precise chromosomal location of the polynucleotides can also be achieved using

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hybridization during chromosomal mapping marking multiple sites and/or multiple chromosomes). Preferred polynucleotides mark a single chromosome or a single site on that chromosome) or in panels (for more likely conserved within gene families, thus increasing the chance of cross correspond to the noncoding regions of the cDNAs because the coding sequences are For chromosome mapping, the polynucleotides can be used individually (to

physical position of the polynucleotide can be used in linkage analysis. Linkage Once a polynucleotide has been mapped to a precise chromosomal location, the ઝ

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analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. (Disease mapping data are found, for example, in V. McKusick, Mendelian Inheritance in Man (available on line through Johns Hopkins University Welch Medical Library).) Assuming I megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential causative genes.

Thus, once coinheritance is established, differences in the polynucleotide and the corresponding gene between affected and unaffected individuals can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected individuals, but not in normal individuals, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal individuals is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this polymorphic polypeptide can be used for further linkage analysis.

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Furthermore, increased or decreased expression of the gene in affected individuals as compared to unaffected individuals can be assessed using polynucleotides of the present invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic marker.

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In addition to the foregoing, a polynucleotide can be used to control gene expression through triple helix formation or antisense DNA or RNA. Both methods rely on binding of the polynucleotide to DNA or RNA. For these techniques, preferred polynucleotides are usually 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques are effective in model systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat disease.

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Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the

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present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell.

The polynucleotides are also useful for identifying individuals from minute biological samples. The United States military, for example, is considering the use of restriction fragment length polymorphism (RFLP) for identification of its personnel. In this technique, an individual's genomic DNA is digested with one or more restriction enzymes, and probed on a Southern blot to yield unique bands for identifying personnel. This method does not suffer from the current limitations of "Dog Tags" which can be lost, switched, or stolen, making positive identification difficult. The polynucleotides of the present invention can be used as additional DNA markers for

The polynucleotides of the present invention can also be used as an alternative to RFLP, by determining the actual base-by-base DNA sequence of selected portions of an individual's genome. These sequences can be used to prepare PCR primers for amplifying and isolating such selected DNA, which can then be sequenced. Using this technique, individuals can be identified because each individual will have a unique set of DNA sequences. Once an unique ID database is established for an individual, positive identification of that individual, living or dead, can be made from extremely small tissue samples.

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Forensic biology also benefits from using DNA-based identification techniques as disclosed herein. DNA sequences taken from very small biological samples such as tissues, e.g., hair or skin, or body fluids, e.g., blood, saliva, semen, etc., can be amplified using PCR. In one prior art technique, gene sequences amplified from polymorphic loci, such as DQa class II HLA gene, are used in forensic biology to identify individuals. (Erlich, H., PCR Technology, Freeman and Co. (1992).) Once these specific polymorphic loci are amplified, they are digested with one or more restriction enzymes, yielding an identifying set of bands on a Southern blot probed with DNA corresponding to the DQa class II HLA gene. Similarly, polynucleotides of the present invention can be used as polymorphic markers for forensic purposes.

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There is also a need for reagents capable of identifying the source of a particular tissue. Such need arises, for example, in forensics when presented with tissue of unknown origin. Appropriate reagents can comprise, for example, DNA probes or primers specific to particular tissue prepared from the sequences of the present invention. Panels of such reagents can identify tissue by species and/or by organ type. In a similar fashion, these reagents can be used to screen tissue cultures for

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In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

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### Uses of the Polypeptides

Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

A polypoptide of the present invention can be used to assay protein levels in a

A polypeptide of the present invention can be used to assay protein levels in a biological sample using antibody-based techniques. For example, protein expression in tissues can be studied with classical immunohistological methods. (Jalkanen, M., et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell. Biol. 105:3087-

20 technetium (99mTc), and fluorescent labels, such as fluorescein and rhodamine, and hiorin.

In addition to assaying secreted protein levels in a biological sample, proteins can also be detected in vivo by imaging. Antibody labels or markers for in vivo imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

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A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 1311, 112In, 99mTc), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously, or intraperitoneally) into the mammal. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, for a human subject, the quantity of radioactivity injected will normally range from about 5 to 20

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millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially accumulate at the location of cells which contain the specific protein. In vivo tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Tumor Imaging: The Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982).)

Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression of a polypeptide of the present invention in cells or body fluid of an individual; (b) comparing the level of gene expression with a standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a disorder.

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Moreover, polypeptides of the present invention can be used to treat disease. For example, patients can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide (e.g., insulin), to supplement absent or decreased levels of a different polypeptide (e.g., hemoglobin S for hemoglobin B), to inhibit the activity of a polypeptide (e.g., an oncogene), to activate the activity of a polypeptide (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor by competing with it for free ligand (e.g., soluble TNF receptors used in reducing inflammation), or to bring about a desired response (e.g., blood vessel growth).

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Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat disease. For example, administration of an antibody directed to a polypeptide of the present invention can bind and reduce overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

At the very least, the polypeptides of the present invention could be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the following biological activities.

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#### <u>ological Activities</u>

35 The polynucleotides and polypeptides of the present invention can be used in assays to test for one or more biological activities. If these polynucleotides and polypeptides do exhibit activity in a particular assay, it is likely that these molecules

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may be involved in the diseases associated with the biological activity. Thus, the polynucleotides and polypeptides could be used to treat the associated disease.

#### mmune Activity

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A polypeptide or polynucleotide of the present invention may be useful in treating deficiencies or disorders of the immune system, by activating or inhibiting the proliferation, differentiation, or mobilization (chemotaxis) of immune cells. Immune cells develop through a process called hematopoiesis, producing myeloid (platelets, red blood cells, neutrophils, and macrophages) and lymphoid (B and T lymphocytes) cells from pluripotent stem cells. The etiology of these immune deficiencies or disorders may be genetic, somatic, such as cancer or some autoimmune disorders, acquired (e.g., by chemotherapy or toxins), or infectious. Moreover, a polynucleotide or polypeptide of the present invention can be used as a marker or detector of a particular immune system disease or disorder.

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treating or detecting deficiencies or disorders of hematopoietic cells. A polypeptide or polynucleotide of the present invention could be used to increase differentiation and proliferation of hematopoietic cells, including the pluripotent stem cells, in an effort to treat those disorders associated with a decrease in certain (or many) types hematopoietic cells. Examples of immunologic deficiency syndromes include, but are not limited to: blood protein disorders (e.g. agammaglobulinemia, dysgammaglobulinemia), ataxia telangiectasia, common variable immunodeficiency, Digeorge Syndrome, HIV infection, HTLV-BLV infection, leukocyte adhesion deficiency syndrome, lymphopenia, phagocyte bactericidal dysfunction, severe combined immunodeficiency (SCIDs), Wiskott-Aldrich Disorder, anemia, thrombocytopenia, or hemoglobinuria.

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Moreover, a polypeptide or polynucleotide of the present invention could also be used to modulate hemostatic (the stopping of bleeding) or thrombolytic activity (clot formation). For example, by increasing hemostatic or thrombolytic activity, a polynucleotide or polypeptide of the present invention could be used to treat blood coagulation disorders (e.g., afibrinogenemia, factor deficiencies), blood platelet disorders (e.g. thrombocytopenia), or wounds resulting from trauma, surgery, or other causes. Alternatively, a polynucleotide or polypeptide of the present invention that can decrease hemostatic or thrombolytic activity could be used to inhibit or dissolve clotting. These molecules could be important in the treatment of heart attacks (infarction), strokes, or scarring.

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A polynucleotide or polypeptide of the present invention may also be useful in treating or detecting autoimmune disorders. Many autoimmune disorders result from

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inappropriate recognition of self as foreign material by immune cells. This inappropriate recognition results in an immune response leading to the destruction of the host tissue. Therefore, the administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-cells, may be an effective therapy in preventing

Examples of autoimmune disorders that can be treated or detected by the present invention include, but are not limited to: Addison's Disease, hemolytic anemia, antiphospholipid syndrome, rheumatoid arthritis, dermatitis, allergic encephalomyelitis, glomerulonephritis, Goodpasture's Syndrome, Graves' Disease, Multiple Sclerosis, Myasthenia Gravis, Neuritis, Ophthalmia, Bullous Pemphigoid, Pemphigus, Polyendocrinopathies, Purpura, Reiter's Disease, Stiff-Man Syndrome, Autoimmune Thyroiditis, Systemic Lupus Erythematosus, Autoimmune Pulmonary Inflammation, Guillain-Barre Syndrome, insulin dependent diabetes mellitis, and autoimmune inflammatory eye disease.

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Similarly, allergic reactions and conditions, such as asthma (particularly allergic asthma) or other respiratory problems, may also be treated by a polypeptide or polynucleotide of the present invention. Moreover, these molecules can be used to treat anaphylaxis, hypersensitivity to an antigenic molecule, or blood group incompatibility.

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A polynucleotide or polypeptide of the present invention may also be used to treat and/or prevent organ rejection or graft-versus-host disease (GVHD). Organ rejection occurs by host immune cell destruction of the transplanted tissue through an immune response. Similarly, an immune response is also involved in GVHD, but, in this case, the foreign transplanted immune cells destroy the host tissues. The administration of a polypeptide or polynucleotide of the present invention that inhibits an immune response, particularly the proliferation, differentiation, or chemotaxis of T-

cells, may be an effective therapy in preventing organ rejection or GVHD.

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Similarly, a polypeptide or polynucleotide of the present invention may also be used to modulate inflammation. For example, the polypeptide or polynucleotide may inhibit the proliferation and differentiation of cells involved in an inflammatory response. These molecules can be used to treat inflammatory conditions, both chronic and acute conditions, including inflammation associated with infection (e.g., septic shock, sepsis, or systemic inflammatory response syndrome (SIRS)), ischemiareperfusion injury, endotoxin lethality, arthritis, complement-mediated hyperacute rejection, nephritis, cytokine or chemokine induced lung injury, inflammatory bowel disease, Crohn's disease, or resulting from over production of cytokines (e.g., TNF or

Hyperproliferative Disorders

A polypeptide or polynucleotide can be used to treat or detect hyperproliferative disorders, including neoplasms. A polypeptide or polynucleotide of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, a polypeptide or polynucleotide of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

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For example, by increasing an immune response, particularly increasing antigenic qualities of the hyperproliferative disorder or by proliferating, differentiating, or mobilizing T-cells, hyperproliferative disorders can be treated. This immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, decreasing an immune response may also be a method of treating hyperproliferative disorders, such as a chemotherapeutic agent.

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Examples of hyperproliferative disorders that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but are not limited to neoplasms located in the: abdomen, bone, breast, digestive system, liver, pancreas, peritoneum, endocrine glands (adrenal, parathyroid, pituitary, testicles, ovary, thymus, thyroid), eye, head and neck, nervous (central and peripheral), lymphatic system, pelvic, skin, soft tissue, spleen, thoracic, and urogenital.

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Similarly, other hyperproliferative disorders can also be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of such hyperproliferative disorders include, but are not limited to: hypergammaglobulinemia, lymphoproliferative disorders, paraproteinemias, purpura, sarcoidosis, Sezary Syndrome, Waldenstron's Macroglobulinemia, Gaucher's Disease, histiocytosis, and

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any other hyperproliferative disease, besides neoplasia, located in an organ system

### Infectious Disease

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A polypeptide or polynucleotide of the present invention can be used to treat or detect infectious agents. For example, by increasing the immune response, particularly increasing the proliferation and differentiation of B and/or T cells, infectious diseases may be treated. The immune response may be increased by either enhancing an existing immune response, or by initiating a new immune response. Alternatively, the polypeptide or polynucleotide of the present invention may also directly inhibit the infectious agent, without necessarily eliciting an immune response.

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Viruses are one example of an infectious agent that can cause disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention. Examples of viruses, include, but are not limited to the following DNA and RNA viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus,

- 5 Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Flaviviridae
  Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes
  Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus,
  Rhabdoviridae), Orthomyxoviridae (e.g., Influenza), Papovaviridae, Parvoviridae,
  Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g.,
- Rotavirus), Retroviridae (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). Viruses falling within these families can cause a variety of diseases or symptoms, including, but not limited to: arthritis, bronchiollitis, encephalitis, eye infections (e.g., conjunctivitis, keratitis), chronic fatigue syndrome, hepatitis (A, B, C, E, Chronic Active, Delta), meningitis, opportunistic infections (e.g., AIDS),
- 15 pneumonia, Burkitt's Lymphoma, chickenpox, hemorrhagic fever, Measles, Mumps, Parainfluenza, Rabies, the common cold, Polio, leukemia, Rubella, sexually transmitted diseases, skin diseases (e.g., Kaposi's, warts), and viremia. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.
- 20 Similarly, bacterial or fungal agents that can cause disease or symptoms and that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following Gram-Negative and Gram-positive bacterial families and fungi: Actinomycetales (e.g., Corynebacterium, Mycobacterium, Norcardia), Aspergillosis, Bacillaceae (e.g., Anthrax, Clostridium), Bacteroidaceae,
- 25 Blastomycosis, Bordetella, Borrelia, Brucellosis, Candidiasis, Campylobacter, Coccidioidomycosis, Cryptococcosis, Dermatocycoses, Enterobacteriaceae (Klebsiella, Salmonella, Serratia, Yersinia), Erysipelothrix, Helicobacter, Legionellosis, Leptospirosis, Listeria, Mycoplasmatales, Neisseriaceae (e.g., Acinetobacter, Gonorrhea, Menigococcal), Pasteurellacea Infections (e.g., Actinobacillus,
- Heamophilus, Pasteurella), Pseudomonas, Rickettsiaceae, Chlamydiaceae, Syphilis, and Staphylococcal. These bacterial or fungal families can cause the following diseases or symptoms, including, but not limited to: bacteremia, endocarditis, eye infections (conjunctivitis, tuberculosis, uveitis), gingivitis, opportunistic infections (e.g., AIDS related infections), paronychia, prosthesis-related infections, Reiter's Disease, Tespiratory tract infections, such as Whooping Cough or Empyema, sepsis, Lyme Disease, Cat-Scratch Disease, Dysentery, Paratyphoid Fever, food poisoning, Typhoid, pneumonia, Gonorrhea, meningitis, Chlamydia, Syphilis, Diphtheria,

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Leprosy, Paratuberculosis, Tuberculosis, Lupus, Botulism, gangrene, tetanus, impetigo, Rheumatic Fever, Scarlet Fever, sexually transmitted diseases, skin diseases (e.g., cellulitis, dermatocycoses), toxemia, urinary tract infections, wound infections. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or diseases.

Moreover, parasitic agents causing disease or symptoms that can be treated or detected by a polynucleotide or polypeptide of the present invention include, but not limited to, the following families: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis, Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis, Leishmaniasis, Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas. These parasites can cause a variety of diseases or symptoms, including, but not limited to: Scabies, Trombiculiasis, eye infections, intestinal disease (e.g., dysentery, giardiasis), liver disease, lung disease, opportunistic infections (e.g., AIDS related), Malaria, pregnancy complications, and toxoplasmosis. A polypeptide or polynucleotide of the present invention can be used to treat or detect any of these symptoms or

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Preferably, treatment using a polypeptide or polynucleotide of the present invention could either be by administering an effective amount of a polypeptide to the patient, or by removing cells from the patient, supplying the cells with a polynucleotide of the present invention, and returning the engineered cells to the patient (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen in a vaccine to raise an immune response against infectious disease.

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diseases.

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#### Regeneration

A polynucleotide or polypeptide of the present invention can be used to differentiate, proliferate, and attract cells, leading to the regeneration of tissues. (See, Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, replace, or protect tissue damaged by congenital defects, trauma (wounds, burns, incisions, or ulcers), age, disease (e.g. osteoporosis, osteocarthritis, periodontal disease, liver failure), surgery, including cosmetic plastic surgery, fibrosis, reperfusion injury, or systemic cytokine damage.

Tissues that could be regenerated using the present invention include organs (e.g., pancreas, liver, intestine, kidney, skin, endothelium), muscle (smooth, skeletal or cardiac), vascular (including vascular endothelium), nervous, hematopoietic, and skeletal (bone, cartilage, tendon, and ligament) tissue. Preferably, regeneration occurs without or decreased scarring. Regeneration also may include angiogenesis.

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Moreover, a polynucleotide or polypeptide of the present invention may increase regeneration of tissues difficult to heal. For example, increased tendon/ligament regeneration would quicken recovery time after damage. A polynucleotide or polypeptide of the present invention could also be used prophylactically in an effort to avoid damage. Specific diseases that could be treated include of tendinitis, carpal tunnel syndrome, and other tendon or ligament defects. A further example of tissue regeneration of non-healing wounds includes pressure ulcers, ulcers associated with vascular insufficiency, surgical, and traumatic wounds.

Similarly, nerve and brain tissue could also be regenerated by using a polynucleotide or polypeptide of the present invention to proliferate and differentiate nerve cells. Diseases that could be treated using this method include central and peripheral nervous system diseases, neuropathies, or mechanical and traumatic disorders (e.g., spinal cord disorders, head trauma, cerebrovascular disease, and stoke). Specifically, diseases associated with peripheral nerve injuries, peripheral neuropathy (e.g., resulting from chemotherapy or other medical therapies), localized neuropathies, and central nervous system diseases (e.g., Alzheimer's disease, Parkinson's disease, Huntington's disease, amyotrophic lateral sclerosis, and Shy-Drager syndrome), could all be treated using the polynucleotide or polypeptide of the present invention.

#### Chemotaxi

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A polynucleotide or polypeptide of the present invention may have chemotaxis activity. A chemotaxic molecule attracts or mobilizes cells (e.g., monocytes, fibroblasts, neutrophils, T-cells, mast cells, eosinophils, epithelial and/or endothelial cells) to a particular site in the body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

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A polynucleotide or polypeptide of the present invention may increase chemotaxic activity of particular cells. These chemotactic molecules can then be used to treat inflammation, infection, hyperproliferative disorders, or any immune system disorder by increasing the number of cells targeted to a particular location in the body. For example, chemotaxic molecules can be used to treat wounds and other trauma to tissues by attracting immune cells to the injured location. Chemotactic molecules of the present invention can also attract fibroblasts, which can be used to treat wounds.

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It is also contemplated that a polynucleotide or polypeptide of the present invention may inhibit chemotactic activity. These molecules could also be used to treat

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disorders. Thus, a polynucleotide or polypeptide of the present invention could be used as an inhibitor of chemotaxis.

#### Binding Activity

A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors),or small molecules

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Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural or functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

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Preferably, the screening for these molecules involves producing appropriate cells which express the polypeptide, either as a secreted protein or on the cell membrane. Preferred cells include cells from mammals, yeast, Drosophila, or *E. coli*. Cells expressing the polypeptide (or cell membrane containing the expressed

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polypeptide) are then preferably contacted with a test compound potentially containing

the molecule to observe binding, stimulation, or inhibition of activity of either the

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The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the candidate compound results in a signal generated by binding to the polypeptide.

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polypeptide or the molecule.

Alternatively, the assay can be carried out using cell-free preparations,
polypeptide/molecule affixed to a solid support, chemical libraries, or natural product
mixtures. The assay may also simply comprise the steps of mixing a candidate
compound with a solution containing a polypeptide, measuring polypeptide/molecule
activity or binding, and comparing the polypeptide/molecule activity or binding to a

Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The

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components

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antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

All of these above assays can be used as diagnostic or prognostic markers. The molecules discovered using these assays can be used to treat disease or to bring about a particular result in a patient (e.g., blood vessel growth) by activating or inhibiting the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptide from suitably manipulated cells or tissues.

Therefore, the invention includes a method of identifying compounds which bind to a polypeptide of the invention comprising the steps of: (a) incubating a candidate binding compound with a polypeptide of the invention; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound with a polypeptide of the invention, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

#### Other Activities

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A polypeptide or polynucleotide of the present invention may also increase or decrease the differentiation or proliferation of embryonic stem cells, besides, as discussed above, hematopoietic lineage.

A polypeptide or polynucleotide of the present invention may also be used to modulate mammalian characteristics, such as body height, weight, hair color, eye color, skin, percentage of adipose tissue, pigmentation, size, and shape (e.g., cosmetic surgery). Similarly, a polypeptide or polynucleotide of the present invention may be used to modulate mammalian metabolism affecting catabolism, anabolism, processing, utilization, and storage of energy.

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A polypeptide or polynucleotide of the present invention may be used to change a mammal's mental state or physical state by influencing biorhythms, caricadic rhythms, depression (including depressive disorders), tendency for violence, tolerance for pain, reproductive capabilities (preferably by Activin or Inhibin-like activity),

hormonal or endocrine levels, appetite, libido, memory, stress, or other cognitive qualities.

A polypeptide or polynucleotide of the present invention may also be used as a food additive or preservative, such as to increase or decrease storage capabilities, fat

content, lipid, protein, carbohydrate, vitamins, minerals, cofactors or other nutritional

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### Other Preferred Embodiments

nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of Other preferred embodiments of the claimed invention include an isolated SEQ ID NO:X wherein X is any integer as defined in Table 1.

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positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of Clone Sequence and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

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positions beginning with the nucleotide at about the position of the 5' Nucleotide of the Start Codon and ending with the nucleotide at about the position of the 3' Nucleotide of Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of the Clone Sequence as defined for SEQ ID NO:X in Table 1.

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Nucleotide of the First Amino Acid of the Signal Peptide and ending with the nucleotide at about the position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the Similarly preferred is a nucleic acid molecule wherein said sequence of range of positions beginning with the nucleotide at about the position of the 5' NO:X in Table 1.

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sequence which is at least 95% identical to a sequence of at least about 150 contiguous Also preferred is an isolated nucleic acid molecule comprising a nucleotide nucleotides in the nucleotide sequence of SEQ ID NO:X.

sequence which is at least 95% identical to a sequence of at least about 500 contiguous Further preferred is an isolated nucleic acid molecule comprising a nucleotide nucleotides in the nucleotide sequence of SEQ ID NO:X.

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ID NO:X beginning with the nucleotide at about the position of the 5' Nucleotide of the nucleotide sequence which is at least 95% identical to the nucleotide sequence of SEQ position of the 3' Nucleotide of the Clone Sequence as defined for SEQ ID NO:X in First Amino Acid of the Signal Peptide and ending with the nucleotide at about the A further preferred embodiment is a nucleic acid molecule comprising a Table 1.

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A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NQ:X. 35

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to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or molecule which hybridizes does not hybridize under stringent hybridization conditions stringent hybridization conditions to a nucleic acid molecule, wherein said nucleic acid Also preferred is an isolated nucleic acid molecule which hybridizes under of only T residues.

Also preferred is a composition of matter comprising a DNA molecule which which DNA molecule is contained in the material deposited with the American Type Culture Collection and given the ATCC Deposit Number shown in Table 1. for said comprises a human cDNA clone identified by a cDNA Clone Identifier in Table cDNA Clone Identifier. 2

Clone Identifier in Table 1, which DNA molecule is contained in the deposit given the nucleotides in the nucleotide sequence of a human cDNA clone identified by a cDNA Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous ATCC Deposit Number shown in Table 1. Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of the complete open reading frame sequence encoded by said human cDNA clone.

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Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone.

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contiguous nucleotides in the nucleotide sequence encoded by said human cDNA clone. A further preferred embodiment is an isolated nucleic acid molecule comprising A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500

A further preferred embodiment is a method for detecting in a biological sample a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by said human cDNA clone. 25

a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer dentified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the nolecule in said sample with a sequence selected from said group and determining to a sequence of at least 50 contiguous nucleotides in a sequence selected from the as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone comprises a step of comparing a nucleotide sequence of at least one nucleic acid ATCC Deposit Number shown for said cDNA clone in Table 1; which method 8 35

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whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

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A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

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Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a secreted protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

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Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the amino acid sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1.

Also preferred is a polypeptide, wherein said sequence of contiguous amino

acids is included in the amino acid sequence of SEQ ID NO:Y in the range of positions beginning with the residue at about the position of the First Amino Acid of the Secreted Portion and ending with the residue at about the Last Amino Acid of the Open Reading Frame as set forth for SEQ ID NO:Y in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

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Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y.

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Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

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Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a secreted portion of the secreted protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 30 contiguous amino acids in the

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clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with arnino acid sequence of the secreted portion of the protein encoded by a human cDNA the ATCC Deposit Number shown for said cDNA clone in Table 1.

clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with Also preferred is an isolated polypeptide comprising an amino acid sequence at arnino acid sequence of the secreted portion of the protein encoded by a human cDNA least 95% identical to a sequence of at least about 100 contiguous amino acids in the the ATCC Deposit Number shown for said cDNA clone in Table 1.

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contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and Also preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the amino acid sequence of the secreted portion of the protein

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human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in sequence of at least 10 contiguous amino acids in a sequence selected from the group the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a polypeptide comprising an arnino acid sequence that is at least 90% identical to a Further preferred is an isolated antibody which binds specifically to a 2

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least 10 contiguous amino acids in a sequence selected from the group consisting of: an Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the and a complete amino acid sequence of a protein encoded by a human cDNA clone molecule in said sample with a sequence selected from said group and determining comprises a step of comparing an amino acid sequence of at least one polypeptide whether the sequence of said polypeptide molecule in said sample is at least 90% ATCC Deposit Number shown for said cDNA clone in Table 1; which method identical to said sequence of at least 10 contiguous amino acids.

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comprising an arnino acid sequence that is at least 90% identical to a sequence of at least Also preferred is the above method wherein said step of comparing an amino polypeptides in said sample to an antibody which binds specifically to a polypeptide 10 contiguous amino acids in a sequence selected from the group consisting of: an acid sequence of at least one polypeptide molecule in said sample with a sequence selected from said group comprises determining the extent of specific binding of 35

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amino acid sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1; identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the and a complete amino acid sequence of a protein encoded by a human cDNA clone ATCC Deposit Number shown for said cDNA clone in Table 1. Also preferred is the above method wherein said step of comparing sequences is performed by comparing the amino acid sequence determined from a polypeptide molecule in said sample with said sequence selected from said group.

by a human cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained biological sample which method comprises a step of detecting polypeptide molecules in said sample, if any, comprising an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. defined in Table 1; and a complete amino acid sequence of a secreted protein encoded Also preferred is a method for identifying the species, tissue or cell type of a consisting of: an amino acid sequence of SEQ ID NO: Y wherein Y is any integer as

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Also preferred is the above method for identifying the species, tissue or cell type sequence of at least 10 contiguous amino acids in a sequence selected from the above sequences, wherein at least one sequence in said panel is at least 90% identical to a molecules comprising an amino acid sequence in a panel of at least two amino acid of a biological sample, which method comprises a step of detecting polypeptide

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obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two arnino acid sequences, wherein at least one sequence in said panel identified in Table 1, which method comprises a step of detecting in a biological sample Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number associated with abnormal structure or expression of a gene encoding a secreted protein Also preferred is a method for diagnosing in a subject a pathological condition sequence of a secreted protein encoded by a human cDNA clone identified by a cDNA sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid is at least 90% identical to a sequence of at least 10 contiguous amino acids in a shown for said cDNA clone in Table 1.

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In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

polypeptide wherein said polypeptide comprises an amino acid sequence that is at least Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding

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90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected 199

protein encoded by a human cDNA clone identified by a cDNA Clone Identifier in Table clone in Table 1. 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA any integer as defined in Table 1; and a complete amino acid sequence of a secreted from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is

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in a prokaryotic host. sequence encoding a polypeptide has been optimized for expression of said polypeptide Also preferred is an isolated nucleic acid molecule, wherein said nucleotide

5 2 ATCC Deposit Number shown for said cDNA clone in Table 1. identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the complete amino acid sequence of a secreted protein encoded by a human cDNA clone sequence of SEQ ID NO: Y wherein Y is any integer as defined in Table 1; and a comprises an amino acid sequence selected from the group consisting of: an amino acid Also preferred is an isolated nucleic acid molecule, wherein said polypeptide

recombinant host cell produced by this method. a recombinant host cell comprising introducing the vector into a host cell, as well as the the recombinant vector produced by this method. Also preferred is a method of making inserting any of the above isolated nucleic acid molecule into a vector. Also preferred is Further preferred is a method of making a recombinant vector comprising

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cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the and an amino acid sequence of a secreted portion of a protein encoded by a human of the First Amino Acid of the Secreted Portion of SEQ ID NO:Y is defined in Table 1; NO:Y beginning with the residue at the position of the First Amino Acid of the Secreted sequence selected from the group consisting of: an amino acid sequence of SEQ ID polypeptide is a secreted portion of a human secreted protein comprising an amino acid expressed and recovering said polypeptide. Also preferred is this method of making an culturing this recombinant host cell under conditions such that said polypeptide is Portion of SEQ ID NO: Y wherein Y is an integer set forth in Table 1 and said position isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said Also preferred is a method of making an isolated polypeptide comprising

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individual a pharmaceutical composition comprising an amount of an isolated level of a secreted protein activity, which method comprises administering to such an Also preferred is a method of treatment of an individual in need of an increased deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The

isolated polypeptide produced by this method is also preferred.

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the level of said protein activity in said individual polypeptide, polynucleotide, or antibody of the claimed invention effective to increase

understood by reference to the following examples, which are provided by way of Having generally described the invention, the same will be more readily

Ś illustration and are not intended as limiting.

# Example 1: Isolation of a Selected cDNA Clone From the Deposited

Each cDNA clone in a cited ATCC deposit is contained in a plasmid vector.

5 5 example, where a particular clone is identified in Table 1 as being isolated in the vector Table 1 identifies the vectors used to construct the cDNA library from which each clone was isolated. In many cases, the vector used to construct the library is a phage vector related plasmid for each phage vector used in constructing the cDNA library. For from which a plasmid has been excised. The table immediately below correlates the

"Lambda Zap," the corresponding deposited clone is in "pBluescript." Vector Used to Construct Library Corresponding Deposited Plasmid

pSport1 pCMVSport 3.0 pCMVSport 2.0 lafmid BA Zap Express pSport1 plafmid BA pBK pCMVSport 3.0 pCMVSport 2.0

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Uni-Zap XR Lambda Zap

pBluescript (pBS) pBluescript (pBS)

pCR®2.1

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5,128,256 and 5,286,636), pBluescript (pBS) (Short, J. M. et al., Nucleic Acids Res XR (U.S. Patent Nos. 5,128, 256 and 5,286,636), Zap Express (U.S. Patent Nos. 16:7583-7600 (1988); Alting-Mees, M. A. and Short, J. M., Nucleic Acids Res. Vectors Lambda Zap (U.S. Patent Nos. 5,128,256 and 5,286,636), Uni-Zap

- မ contains a neomycin resistance gene. Both can be transformed into E. coli strain XL-1 Road, La Jolla, CA, 92037. pBS contains an ampicillin resistance gene and pBK commercially available from Stratagene Cloning Systems, Inc., 11011 N. Torrey Pines 17:9494 (1989)) and pBK (Alting-Mees, M. A. et al., Strategies 5:58-61 (1992)) are Blue, also available from Stratagene. pBS comes in 4 forms SK+, SK-, KS+ and KS
- ઝ are the first sites on each respective end of the linker). "+" or "-" refer to the orientation sequences which flank the polylinker region ("S" is for SacI and "K" is for KpnI which The S and K refers to the orientation of the polylinker to the T7 and T3 primer

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of the f1 origin of replication ("ori"), such that in one orientation, single stranded rescue initialed from the f1 ori generates sense strand DNA and in the other, antisense.

Vectors pSport1, pCMVSport 2.0 and pCMVSport 3.0, were obtained from Life Technologies, Inc., P. O. Box 6009, Gaithersburg, MD 20897. All Sport vectors contain an ampicillin resistance gene and may be transformed into E. coli strain DH10B, also available from Life Technologies. (See, for instance, Gruber, C. E., et

al., Focus 15:59 (1993).) Vector lafmid BA (Bento Soares, Columbia University, NY)

contains an ampicillin resistance gene and can be transformed into E. coli strain XL-1

Blue. Vector pCR®2.1, which is available from Invitrogen, 1600 Faraday Avenue,
10. Carlsbad, CA 92008, contains an ampicillin resistance gene and may be transformed
into E. coli strain DH10B, available from Life Technologies. (See, for instance, Clark,
1. M., Nuc. Acids Res. 16:9677-9686 (1988) and Mead, D. et al., Bio/Technology 9:
(1991).) Preferably, a polynucleotide of the present invention does not comprise the
phage vector sequences identified for the particular clone in Table 1, as well as the
corresponding plasmid vector sequences designated above.

The deposited material in the sample assigned the ATCC Deposit Number cited in Table 1 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing the same ATCC Deposit Number contain at least a plasmid for each cDNA clone identified in Table 1. Typically, each ATCC deposit sample cited in Table 1 comprises a mixture of approximately equal amounts (by weight) of about 50 plasmid DNAs, each containing a different cDNA clone; but such a deposit sample may include plasmids for more or less than 50 cDNA clones, up to about 500 cDNA clones.

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Two approaches can be used to isolate a particular clone from the deposited sample of plasmid DNAs cited for that clone in Table 1. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to SEQ ID NO:X.

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Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported.

The oligonucleotide is labeled, for instance, with <sup>13</sup>P-Y-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).) The plasmid mixture is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection

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agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening (e.g., Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1,93 to

5 1.104), or other techniques known to those of skill in the art.

Alternatively, two primers of 17-20 nucleotides derived from both ends of the SEQ ID NO:X (i.e., within the region of SEQ ID NO:X bounded by the 5' NT and the 3' NT of the clone defined in Table 1) are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction

10 is carried out under routine conditions, for instance, in 25 µl of reaction mixkure with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl<sub>1</sub>, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dCTP, dTTP; 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR((denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are

15 performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product. Several methods are available for the identification of the 5' or 3' non-coding portions of a gene which may not be present in the deposited clone. These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols which are well known in the art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., Nucleic Acids Res. 21(7):1683-1684 (1993).)

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Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be sequenced and used to generate the full length gene.

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This above method starts with total RNA isolated from the desired source, although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged

prospurate in recessary to contained by prospurate groups on degraded or damaged

35 RNA which may interfere with the later RNA ligase step. The phosphatase should then
be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to

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remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.

This modified RNA preparation is used as a template for first strand cDNA synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene.

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# Example 2: Isolation of Genomic Clones Corresponding to a Polynucleotide

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A human genomic P1 library (Genomic Systems, Inc.) is screened by PCR using primers selected for the cDNA sequence corresponding to SEQ ID NO:X., according to the method described in Example 1. (See also, Sambrook.)

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## Example 3: Tissue Distribution of Polypeptide

Tissue distribution of mRNA expression of polynucleotides of the present invention is determined using protocols for Northern blot analysis, described by, among others, Sambrook et al. For example, a cDNA probe produced by the method described in Example 1 is labeled with P<sup>27</sup> using the rediprime<sup>TM</sup> DNA labeling system (Amersham Life Science), according to manufacturer's instructions. After labeling, the probe is purified using CHROMA SPIN-100<sup>TM</sup> column (Clontech Laboratories, Inc.), according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various human tissues for mRNA expression.

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Multiple Tissue Northern (MTN) blots containing various human tissues (H) or human immune system tissues (IM) (Clontech) are examined with the labeled probe using ExpressHyb<sup>TM</sup> hybridization solution (Clontech) according to manufacturer's protocol number PT1190-1. Following hybridization and washing, the blots are

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30 mounted and exposed to film at -70°C overnight, and the films developed according to standard procedures.

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# Example 4: Chromosomal Mapping of the Polynucleotides

An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This primer set is then used in a polymerase chain reaction under the following set of

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conditions: 30 seconds, 95°C; I minute, 56°C; I minute, 70°C. This cycle is repeated 32 times followed by one 5 minute cycle at 70°C. Human, mouse, and hamster DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions is analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

## Example 5: Bacterial Expression of a Polypeptide

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A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and Xbal, at the 5' end of the primers in order to clone the amplified product into the expression vector. For example, BamHI and Xbal correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (AmpP), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

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The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, which expresses the lacI repressor and also confers kanamycin resistance (Kanf). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis. Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D. 60) of between 0.4 and 0.6. IPTG (Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1 mM.

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Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic

IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased

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gene expression.

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QIAGEN, Inc., supra). Proteins with a 6 x His tag bind to the Ni-NTA resin with high onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from removed by centrifugation, and the supernatant containing the polypeptide is loaded agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4°C. The cell debris is affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., supra).

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Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

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The purified protein is then renatured by dialyzing it against phosphate-buffered recommended conditions are as follows: renature using a linear 6M-1M urea gradient in saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the renaturation the proteins are eluted by the addition of 250 mM immidazole. Immidazole 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer protein can be successfully refolded while immobilized on the Ni-NTA column. The The renaturation should be performed over a period of 1.5 hours or more. After plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

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In addition to the above expression vector, the present invention further includes a selection marker, 2) an E. coli origin of replication, 3) a T5 phage promoter sequence, 4) two lac operator sequences, 5) a Shine-Delgamo sequence, and 6) the lactose operon Number XXXXXX.) This vector contains: 1) a neomycinphosphotransferase gene as inked to a polynucleotide of the present invention, called pHE4a. (ATCC Accession repressor gene (lacIq). The origin of replication (oriC) is derived from pUC19 (LTI, an expression vector comprising phage operator and promoter elements operatively Gaithersburg, MD). The promoter sequence and operator sequences are made synthetically. 2 23

Xbal, BamHI, Xhol, or Asp718, running the restricted product on a gel, and isolating insert is generated according to the PCR protocol described in Example 1, using PCR he larger fragment (the stuffer fragment should be about 310 base pairs). The DNA DNA can be inserted into the pHEa by restricting the vector with Ndel and Asp718 (3' primer). The PCR insert is gel purified and restricted with compatible primers having restriction sites for Ndel (5' primer) and Xbal, BamHI, Xhol, or enzymes. The insert and vector are ligated according to standard protocols. 9

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The engineered vector could easily be substituted in the above protocol to express protein in a bacterial system.

# Example 6: Purification of a Polypeptide from an Inclusion Body

in E coli when it is present in the form of inclusion bodies. Unless otherwise specified, The following alternative method can be used to purify a polypeptide expressed all of the following steps are conducted at 4-10°C.

Jpon completion of the production phase of the  $\it E.~coli$  fermentation, the cell culture is cooled to 4-10°C and the cells harvested by continuous centrifugation at

- 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer. 2
- (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by The cells are then lysed by passing the solution through a microfluidizer NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4. 2
- 13 ndrochloride (GuHCI) for 2-4 hours. After 7000 xg centrifugation for 15 min., the The resulting washed inclusion bodies are solubilized with 1.5 M guanidine pellet is discarded and the polypeptide containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

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the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 Following high speed centrifugation (30,000 xg) to remove insoluble particles, volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps. 22

filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a

Fractions are collected and further analyzed by SDS-PAGE. stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored

columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated of water. The diluted sample is then loaded onto a previously prepared set of tandem volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 Fractions containing the polypeptide are then pooled and mixed with 4 volumes

5 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A<sub>280</sub> instance, by 16% SDS-PAGE) are then pooled. monitoring of the effluent. Fractions containing the polypeptide (determined, for

refolding and purification steps. No major contaminant bands should be observed from The resultant polypeptide should exhibit greater than 95% purity after the above

2 Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. the LPS content is less than 0.1 ng/ml according to LAL assays. The purified protein can also be tested for endotoxin/LPS contamination, and typically

### Expression System Example 7: Cloning and Expression of a Polypeptide in a Baculovirus

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polyhedrin promoter of the Autographa californica nuclear polyhedrosis virus into a baculovirus to express a polypeptide. This expression vector contains the strong (AcMNPV) followed by convenient restriction sites such as BamHI, Xba I and In this example, the plasmid shuttle vector pA2 is used to insert a polynucleotide

- 25 same orientation, followed by the polyadenylation signal of the polyhedrin gene. The polyadenylation. For easy selection of recombinant virus, the plasmid contains the Asp718. The polyadenylation site of the simian virus 40 ("SV40") is used for efficient inserted genes are flanked on both sides by viral sequences for cell-mediated beta-galactosidase gene from E. coli under control of a weak Drosophila promoter in the
- 30 express the cloned polynucleotide. homologous recombination with wild-type viral DNA to generate a viable virus that

translation, secretion and the like, including a signal peptide and an in-frame AUG as long as the construct provides appropriately located signals for transcription as pAc373, pVL941, and pAcIM1, as one skilled in the art would readily appreciate, as Many other baculovirus vectors can be used in place of the vector above, such

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required. Such vectors are described, for instance, in Luckow et al., Virology 170:31-Specifically, the cDNA sequence contained in the deposited clone, including the

is amplified using the PCR protocol described in Example 1. If the naturally occurring second signal peptide. Alternatively, the vector can be modified (pA2 GP) to include a signal sequence is used to produce the secreted protein, the pA2 vector does not need a AUG initiation codon and the naturally associated leader sequence identified in Table 1 baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures,"

5 Texas Agricultural Experimental Station Bulletin No. 1555 (1987).

with appropriate restriction enzymes and again purified on a 1% agarose gel. available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested The amplified fragment is isolated from a 1% agarose gel using a commercially

2 optionally, can be dephosphorylated using calf intestinal phosphatase, using routine commercially available kit ("Geneclean" BIO 101 Inc., La Jolla, Ca.). procedures known in the art. The DNA is then isolated from a 1% agarose gel using a The plasmid is digested with the corresponding restriction enzymes and

DNA ligase. E. coli HB101 or other suitable E. coli hosts such as XL-1 Blue The fragment and the dephosphorylated plasmid are ligated together with T4

- 20 (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation electrophoresis. The sequence of the cloned fragment is confirmed by DNA digesting DNA from individual colonies and analyzing the digestion product by gel mixture and spread on culture plates. Bacteria containing the plasmid are identified by
- 25 of a commercially available linearized baculovirus DNA ("BaculoGold™ baculovirus Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One μg of DNA", Pharmingen, San Diego, CA), using the lipofection method described by BaculoGold<sup>TM</sup> virus DNA and 5  $\mu g$  of the plasmid are mixed in a sterile well of a Five  $\mu g$  of a plasmid containing the polynucleotide is co-transfected with 1.0  $\mu g$
- ဗ microtiter plate containing 50 µl of serum-free Grace's medium (Life Technologies incubated for 5 hours at 27° C. The transfection solution is then removed from the plate added, mixed and incubated for 15 minutes at room temperature. Then the transfection Inc., Gaithersburg, MD). Afterwards, 10 µl Lipofectin plus 90 µl Grace's medium are and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. tissue culture plate with 1 ml Grace's medium without serum. The plate is then mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm
- ઝ Cultivation is then continued at 27° C for four days

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After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, supra. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing the recombinant baculovirus is used to infect Sf9 cells seeded in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4° C.

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To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 µCi of <sup>35</sup>S-methionine and 5 µCi <sup>35</sup>S-cysteine (available from Amersham) are added. The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

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Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the produced protein.

# 25 Example 8: Expression of a Polypeptide in Mammalian Cells

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which mediates the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLVI, HIVI and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

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35 Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden),

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pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSport 2.0, and pCMVSport 3.0. Mammalian host cells that could be used include, human Hela, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO)

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Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transfection with a selectable marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transfected cells.

The transfected gene can also be amplified to express large amounts of the encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Fage, M. J. and Sydenham, M. A., Biotechnology 9:64-68 (1991).) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., Biochem J. 227:277-279 (1991); Bebbington et al., Bio/Technology 10:169-175 (1992). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated intola

chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the

production of proteins.

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Derivatives of the plasmid pSV2-dhfr (ATCC Accession No. 37146), the expression vectors pC4 (ATCC Accession No. 209646) and pC6 (ATCC Accession No. 209647) contain the strong promoter (LTR) of the Rous Sarcoma Virus (Cullen et al., Molecular and Cellular Biology, 438-447 (March, 1985)) plus a fragment of the CMV-enhancer (Boshart et al., Cell 41:521-530 (1985). Multiple cloning sites, e.g., with the restriction enzyme cleavage sites BamHI, Xbal and Asp718, facilitate the cloning of the gene of interest. The vectors also contain the 3' intron, the polyadenylation and termination signal of the rat preproinsulin gene, and the mouse DHFR gene under control of the SV40 early promoter.

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Specifically, the plasmid pC6, for example, is digested with appropriate restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from a 1% agarose gel.

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A polynucleotide of the present invention is amplified according to the protocol 35 outlined in Example 1. If the naturally occurring signal sequence is used to produce the secreted protein, the vector does not need a second signal peptide. Alternatively, if the

naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.)

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("Geneclean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

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The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coll* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

10 pC6 using, for instance, restriction enzyme analysis.
Chinese hamster ovary cells lacking an active DHFR gene is used for transfection. Five μg of the expression plasmid pC6 is cotransfected with 0.5 μg of the plasmid pSVneo using lipofectin (Felgner et al., supra). The plasmid pSV2-neo

contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that

alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of metothrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri

20 dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μM, 2 μM, 5 μM, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 -

25 200 μM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

### Example 9: Protein Fusions

The polypeptides of the present invention are preferably fused to other proteins.

These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG domains, and maltose binding protein facilitates purification. (See Example 5; see also EP A 394,827;

Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and albumin increases the halfilfe time in vivo. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having

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more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule, or the protocol described in Example 5.

Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate cloning into an expression vector, preferably a mammalian expression vector.

For example, if pC4 (Accession No.209646) is used, the human Fc portion can be ligated into the BamHI cloning site. Note that the 3' BamHI site should be destroyed. Next, the vector containing the human Fc portion is re-restricted with BamHI, linearizing the vector, and a polynucleotide of the present invention, isolated by the PCR protocol described in Example 1, is ligated into this BamHI site. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

If the naturally occurring signal sequence is used to produce the secreted protein, pC4 does not need a second signal peptide. Alternatively, if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (Sec. e.g., WO 96/34891.)

Human IgG Fc region:

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23 35 છ GACCTGCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGTGGAGTGGGA ATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCCGAGAACCACAGGT GCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTACAAC CAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTCACATGCGTGGTGGT GGGATCCGGAGCCCAAATCTTCTGACAAAACTCACACATGCCCACCGTGCC GACGGCCGCGACTCTAGAGGAT (SEQ ID NO:1) ACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGGGTAAATGAGTGC GGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGC ACTCCGACGGCTCCTTCTTCCTCTACAGCAAGCTCACCGTGGACAAGAGCA GAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGCTGG GTACACCCTGCCCCATCCCGGGATGAGCTGACCAAGAACCAGGTCAGCCT AATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCAACCCCC AGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACTGGCTG GGACGTAAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGGACG CAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCCTCTTCCCCCCAAAACC

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# Example 10: Production of an Antibody from a Polypeptide

The antibodies of the present invention can be prepared by a variety of methods. containing polyclonal antibodies. In a preferred method, a preparation of the secreted protein is prepared and purified to render it substantially free of natural contaminants. (See, Current Protocols, Chapter 2.) For example, cells expressing a polypeptide of Such a preparation is then introduced into an animal in order to produce polyclonal the present invention is administered to an animal to induce the production of sera antisera of greater specific activity.

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preferably, with a secreted polypeptide-expressing cell. Such cells may be cultured in iny suitable tissue culture medium; however, it is preferable to culture cells in Earle's 256:495 (1975); Köhler et al., Eur. J. Immunol. 6:511 (1976); Köhler et al., Eur. J. about 56°C), and supplemented with about 10 g/l of nonessential amino acids, about modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at Immunol. 6:292 (1976); Hammerling et al., in: Monoclonal Antibodies and T-Cell Hybridomas, Elsevier, N.Y., pp. 563-681 (1981).) In general, such procedures monoclonal antibodies (or protein binding fragments thereof). Such monoclonal In the most preferred method, the antibodies of the present invention are antibodies can be prepared using hybridoma technology. (Köhler et al., Nature involve immunizing an animal (preferably a mouse) with polypeptide or, more 1,000 U/ml of penicillin, and about 100 µg/ml of streptomycin. 13 2 20

described by Wands et al. (Gastroenterology 80:225-232 (1981).) The hybridoma cells The splenocytes of such mice are extracted and fused with a suitable myeloma (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are obtained through such a selection are then assayed to identify clones which secrete cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line selectively maintained in HAT medium, and then cloned by limiting dilution as antibodies capable of binding the polypeptide. 23

Alternatively, additional antibodies capable of binding to the polypeptide can be possible to obtain an antibody which binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is 35 8

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whose ability to bind to the protein-specific antibody can be blocked by the polypeptide. Such antibodies comprise anti-idiotypic antibodies to the protein-specific antibody and can be used to immunize an animal to induce formation of further protein-specific

fragments are typically produced by proteolytic cleavage, using enzymes such as papain It will be appreciated that Fab and F(ab')2 and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). Alternatively, secreted protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry. S 9

'humanized" chimeric monoclonal antibodies. Such antibodies can be produced using genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art. For in vivo use of antibodies in humans, it may be preferable to use

(See, for review, Morrison, Science 229.1202 (1985); Oi et al., BioTechniques 4.214 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP:171496; (1985).) 12

# Example 11: Production Of Secreted Protein For High-Throughput

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The following protocol produces a supernatant containing a polypeptide to be tested. This supernatant can then be used in the Screening Assays described in Examples 13-20.

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(note: a 12-channel pipetter may be used with tips on every other channel). Aspirate off (1mg/ml in PBS) 1:20 in PBS (w/o calcium or magnesium 17-516F Biowhittaker) for a working solution of 50ug/ml. Add 200 ul of this solution to each well (24 well plates) the Poly-D-Lysine solution and rinse with 1ml PBS (Phosphate Buffered Saline). The First, dilute Poly-D-Lysine (644 587 Boehringer-Mannheim) stock solution and incubate at RT for 20 minutes. Be sure to distribute the solution over each well

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DMEM(Dulbecco's Modified Eagle Medium)(with 4.5 G/L glucose and  $\dot{L}_{\!\!-}$  glutamine Plate 293T cells (do not carry cells past P+20) at 2 x 10<sup>3</sup> cells/well in .5ml (12-604F Biowhittaker))/10% heat inactivated FBS(14-503F Biowhittaker)/1x Penstrep(17-602E Biowhittaker). Let the cells grow overnight.

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PBS should remain in the well until just prior to plating the cells and plates may be

poly-lysine coated in advance for up to two weeks.

vector containing a polynucleotide insert, produced by the methods described in With a small volume multi-channel pipetter, aliquot approximately 2ug of an expression (18324-012 Gibco/BRL) and 5ml Optimem I (31985070 Gibco/BRL)/96-well plate. The next day, mix together in a sterile solution basin: 300 ul Lipofectamine

5 S transfections. control, one plate of vector DNA lacking an insert should be transfected with each set of minutes, use a multi-channel pipetter to add 150ul Optimem I to each well. As a Pipette up and down gently to mix. Incubate at RT 15-45 minutes. After about 20 multi-channel pipetter, add 50ul of the Lipofectamine/Optimem I mixture to each well Examples 8 or 9, into an appropriately labeled 96-well round bottom plate. With a

the even wells, to each row on the 24-well plates. Incubate at 37°C for 6 hours. adds the 200ul of DNA/Lipofectamine/Optimem I complex to the odd wells first, then to PBS rinse, and person B, using a12-channel pipetter with tips on every other channel, cells, and then person B rinses each well with .5-1ml PBS. Person A then aspirates off much time on PBS. First, person A aspirates off the media from four 24-well plates of tasks. By tag-teaming, hands on time is cut in half, and the cells do not spend too Preferably, the transfection should be performed by tag-teaming the following

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solution). Filter the media and collect 50 ul for endotoxin assay in 15ml polystyrene (BSA (81-068-3 Bayer) 100gm dissolved in 1L DMEM for a 10% BSA stock with 1x penstrep, or CHO-5 media (see below) with 2mm glutamine and 1x penstrep. While cells are incubating, prepare appropriate media, either 1%BSA in DMEM

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conical

the incubation period. Person A aspirates off the transfection media, while person B The transfection reaction is terminated, preferably by tag-teaming, at the end of

adds 1.5ml appropriate media to each well. Incubate at 37°C for 45 or 72 hours depending on the media used: 1%BSA for 45 hours or CHO-5 for 72 hours. On day four, using a 300ul multichannel pipetter, aliquot 600ul in one 1ml deep

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each well can then be used in the assays described in Examples 13-20 well plate and the remaining supernatant into a 2ml deep well. The supernatants from

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activity in a particular assay. proteins, which are then secreted into the supernatant. Thus, the invention further provides a method of identifying the protein in the supernatant characterized by an directly (e.g., as a secreted protein) or by the polypeptide inducing expression of other described below using a supernatant, the activity originates from either the polypeptide It is specifically understood that when activity is obtained in any of the assays

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## HGS-CHO-5 medium formulation:

### Inorganic Salts

_					_	_					
ZnSO <sub>4</sub> -7H <sub>2</sub> O	Na,HPO4	NaH,PO,-H,0	NaHCO,	NaCl	MgSO,	MgCl <sub>2</sub>	KC!	FeSO,-7H,0	$Fe(NO_3)_3-9H_2O$	CuSO <sub>4</sub> -5H <sub>2</sub> O	CaCiz (annya)
.4320	71.02	62.50	2400.0	6995.50	48.84	28.64	311.80	0.417	0.050	0.00130	116.6 mg/L

#### S Lipids

_	_	_	_	_	_	_		_	_	_		-	
Tween 80	Stearic Acid	Pluronic F-68	Palmitic Acid	Palmitric Acid	Oleic Acid	Myristic Acid	Linolenic Acid	Linoleic Acid	Tocopherol-Acetate	DL-alpha-	Cholesterol	Arachidonic Acid	
2.20	0.010	100	0.010	0.010	0.010	0.010	0.010	0.0520		.070	1.022	.002 mg/L	

### Carbon Source

D-Glucose	Car post control
4551 mg/L	

#### Amino Acids

L- Alanine	130.85 mg/ml
L-Arginine-HCL	147.50
L-Asparagine-H <sub>7</sub> 0	7.50
L-Aspartic Acid	6.65
L-Cystine-2HCL-	29.56
L-Cystine-2HCL	31.29
L-Glutamic Acid	7.35
L-Glutamine	365.0
Glycine	18.75
L-Histidine-HCL-	52.48

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H.0	
L-Isoleucine	106.97
L-Leucine	111.45
L-Lysine HCL	163.75
L-Methionine	32.34
L-Phenylalainine	68.48
L-Proline	40.0
L-Serine	26.25
L-Threonine	101.05
L-Tryptophan	19.22
L-Tryrosine-2Na- 2H,0	91.79
L-Valine	99.65

#### Vitamins

Biotin	0.0035 mg/L
D-Ca Pantothenate	3.24
Choline Chloride	11.78
Folic Acid	4.65
i-Inositol	15.60
Niacinamide	3.02
Pyridoxal HCL	3.00
Pyridoxine HCL	0.031
Riboflavin	0.319
Thiamine HCL	3.17
Thymidine	0.365
Vitamin B <sub>1</sub> ,	0890

### Other Components

HEPES Buffer	25 mM
Na Hypoxanthine	2.39 mg/L
Lipoic Acid	0.105
Sodium Putrescine-2HCL	0.081
Sodium Pyruvate	55.0
Sodium Selenite	0.0067
Ethanolamine	20uM
Ferric Citrate	0.122
Methyl-B-Cyclodextrin complexed with	41.70
Linoleic Acid	
Methyl-B-Cyclodextrin complexed with	33.33
Oleic Acid	
Methyl-B-Cyclodextrin complexed with	01
Retinal Acetate	

Adjust osmolarity to 327 mOsm

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Example 12: Construction of GAS Reporter Construct

responsive element ("ISRE"), located in the promoter of many genes. The binding of a One signal transduction pathway involved in the differentiation and proliferation of cells is called the Jaks-STATs pathway. Activated proteins in the Jaks-STATs pathway bind to gamma activation site "GAS" elements or interferon-sensitive protein to these elements alter the expression of the associated gene.

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GAS and ISRE elements are recognized by a class of transcription factors called higher concentrations in other cells including myeloid cells. It can be activated in tissue many cell types though it has been found in T helper class I, cells after treatment with Stat2 (as response to IFN-alpha is widespread). Stat4 is more restricted and is not in members of the STATs family. Stat1 and Stat3 are present in many cell types, as is IL-12. Stat5 was originally called mammary growth factor, but has been found at Signal Transducers and Activators of Transcription, or "STATs." There are six culture cells by many cytokines. 2

The STATs are activated to translocate from the cytoplasm to the nucleus upon family. Jaks represent a distinct family of soluble tyrosine kinases and include Tyk2, Jak1, Jak2, and Jak3. These kinases display significant sequence similarily and are tyrosine phosphorylation by a set of kinases known as the Janus Kinase ("Jaks") generally catalytically inactive in resting cells. 12

below. (Adapted from review by Schidler and Darnell, Ann. Rev. Biochem. 64:621-51 groups: (a) Class 1 includes receptors for IL-2, IL-3, IL-4, IL-6, IL-7, IL-9, IL-11, IL-12, IL-15, Epo, PRL, GH, G-CSF, GM-CSF, LIF, CNTF, and thrombopoietin; and The Jaks are activated by a wide range of receptors summarized in the Table (1995).) A cytokine receptor family, capable of activating Jaks, is divided into two ຊ

conserved cysteine motif (a set of four conserved cysteines and one tryptophan) and a WSXWS motif (a membrane proxial region encoding Trp-Ser-Xxx-Trp; Ser (SEQ ID (b) Class 2 includes IFN-a, IFN-g, and IL-10. The Class 1 receptors share a 22

activate STATs, which then translocate and bind to GAS elements. This entire process Thus, on binding of a ligand to a receptor, Jaks are activated, which in turn is encompassed in the Jaks-STATs signal transduction pathway. 30

proliferation and differentiation of cells. For example, growth factors and cytokines are known to activate the Jaks-STATs pathway. (See Table below.) Thus, by using GAS Therefore, activation of the Jaks-STATs pathway, reflected by the binding of elements linked to reporter molecules, activators of the Jaks-STATs pathway can be the GAS or the ISRE element, can be used to indicate proteins involved in the identified.

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	To construct a synthetic GAS containing promoter element, which is used in the
	Biological Assays described in Examples 13-14, a PCR based strategy is employed to
	generate a GAS-SV40 promoter sequence. The 5' primer contains four tandem copies
	of the GAS binding site found in the IRF1 promoter and previously demonstrated to
v	bind STATs upon induction with a range of cytokines (Rothman et al., Immunity
	1:457-468 (1994).), although other GAS or ISRE elements can be used instead. The 5'
,	primer also contains 18bp of sequence complementary to the SV40 early promoter
	sequence and is flanked with an XhoI site. The sequence of the 5' primer is:
	5'.GCGCCTCGAGATTTCCCCCGAAATCTAGATTTCCCCCGAAATGATTTCCCCCG
10	10 AAATGATTTCCCCGAAATATCTGCCATCTCAATTAG:3' (SEO ID NO:3)

NO:4) with a Hind III site: 5':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID The downstream primer is complementary to the SV40 promoter and is flanked

the B-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is with forward and reverse primers confirms that the insert contains the following digested with Xhol/Hind III and subcloned into BLSK2-. (Stratagene.) Sequencing sequence: PCR amplification is performed using the SV40 promoter template present in

20 ATTTCCCCGAAATATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCC 5':CICGAGATTTCCCCGAAATCTAGATTTCCCCGAAATGATTTCCCCGAAATG CCCATGGCTGACTAATTTTTTTTATTTATGCAGAGGCCGAGGCCGCCTCGGC TGCAAA<u>AAGCTT</u>:3' (SEQ ID NO:5) CTAACTCCGCCCATCCCGCCCCTAACTCCGCCCAGTTCCGCCCATTCTCCGC

မ be used instead of SEAP include chloramphenicol acetyltransferase (CAT), luciferase, reporter construct is next engineered. Here, the reporter molecule is a secreted alkaline phosphatase, or "SEAP." Clearly, however, any reporter molecule can be instead of detectable by an antibody. alkaline phosphatase, B-galactosidase, green fluorescent protein (GFP), or any protein SEAP, in this or in any of the other Examples. Well known reporter molecules that car With this GAS promoter element linked to the SV40 promoter, a GAS:SEAP2

3

PDGF CSF-1

<u>....</u>

GAS (not IRF1)

1,3

GAS (IRF1)

Receptor Tyrosine Kinases
2

6

EPO CAS>IRF1=IFP>>Ly6)

1,3,5 5

GAS(B-

35

IL-5 (myeloid)
GM-CSF (myeloid) (IRF1>IFP>>Ly6)

GAS GAS

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GAS

Growth hormone family GH

30

gp140 family IL-3 (myeloid)

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L-9 (lymphocytes) 3 (lymphocyte) (lymphocytes)

GAS GAS

L-4 (lymph/mycloid) ->Ly6)(IgH) L-2 (lymphocytes)

1,3,5 6

(IRFI = IFP)

2

20

2-C family

5

II-11(Pleiotrohic)
OnM(Pleiotrohic)
LIF(Pleiotrohic)
CNTF(Pleiotrohic)
G-CSF(Pleiotrohic)
II-12(Pleiotrohic)

5

IFN family IFN-a/B

(IRF1>Lys6>IFP)

ISRE Ligand

tyk2

Jakl Jak2

Jak3

1,2,3

ISRE GAS

1,3

JAKs

STATS

GAS(elements) or

gp130 family IL-6 (Pleiotrohic)

1,3

GAS

IRF1>Lys6>IFP

35 element, to create the GAS-SEAP vector. However, this vector does not contain a Xhol, effectively replacing the SV40 promoter with the amplified GAS:SV40 promoter subcloned into the pSEAP-Promoter vector obtained from Clontech using HindIII and The above sequence confirmed synthetic GAS-SV40 promoter element is

neomycin resistance gene, and therefore, is not preferred for mammalian expression

SEAP reporter, the GAS-SEAP cassette is removed from the GAS-SEAP vector using Sall and Notl, and inserted into a backbone vector containing the neomycin resistance mammalian cells, this vector can then be used as a reporter molecule for GAS binding gene, such as pGFP-1 (Clontech), using these restriction sites in the multiple cloning Thus, in order to generate mammalian stable cell lines expressing the GASsite, to create the GAS-SEAP/Neo vector. Once this vector is transfected into as described in Examples 13-14.

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containing NFK-B and EGR promoter sequences are described in Examples 15 and 16. with a different promoter sequence. For example, construction of reporter molecules Other constructs can be made using the above description and replacing GAS these Examples. For instance, SRE, IL-2, NFAT, or Osteocalcin promoters can be However, many other promoters can be substituted using the protocols described in construct activity, such as HELA (epithelial), HUVEC (endothelial), Reh (B-cell), 2/NFAT, or NF-KB/GAS). Similarly, other cell lines can be used to test reporter substituted, alone or in combination (e.g., GAS/NF-KB/EGR, GAS/NF-KB, II. Saos-2 (osteoblast), HUVAC (aortic), or Cardiomyocyte. 2 13

## Example 13: High-Throughput Screening Assay for T-cell Activity, 8

Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS Accession No. TIB-152), although Molt-3 cells (ATCC Accession No. CRL-1552) and cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. such as growth factors and cytokines, that may proliferate or differentiate T-cells. T-The following protocol is used to assess T-cell activity by identifying factors, signal transduction pathway. The T-cell used in this assay is Jurkat T-cells (ATCC Molt-4 cells (ATCC Accession No. CRL-1582) cells can also be used. 23

20,000 cells per well and transfectants resistant to 1 mg/ml genticin selected. Resistant colonies are expanded and then tested for their response to increasing concentrations of Jurkat T-cells are lymphoblastic CD4+ Th1 helper cells. In order to generate stable cell lines, approximately 2 million Jurkat cells are transfected with the GASdescribed below). The transfected cells are seeded to a density of approximately SEAP/neo vector using DMRIE-C (Life Technologies)(transfection procedure interferon gamma. The dose response of a selected clone is demonstrated.

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generate sufficient cells for multiple 96 well plates. Jurkat cells are maintained in RPMI containing 200 ul of cells. Thus, it is either scaled up, or performed in multiple to Specifically, the following protocol will yield sufficient cells for 75 wells 35

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+ 10% serum with 1%Pen-Strep. Combine 2.5 mls of OPTI-MEM (Life Technologies) with 10 ug of plasmid DNA in a T25 flask. Add 2.5 ml OPTI-MEM containing 50 ul of DMRIE-C and incubate at room temperature for 15-45 mins.

concentration of 10' cells/ml. Then add 1 ml of 1 x 10' cells in OPTI-MEM to T25 flask and incubate at 37°C for 6 hrs. After the incubation, add 10 ml of RPMI + 15% serum. During the incubation period, count cell concentration, spin down the required number of cells (107 per transfection), and resuspend in OPTI-MEM to a final

serum, I mg/ml Genticin, and 1% Pen-Strep. These cells are treated with supernatants The Jurkat: GAS-SEAP stable reporter lines are maintained in RPMI + 10% containing a polypeptide as produced by the protocol described in Example 11.

2

On the day of treatment with the supernatant, the cells should be washed and resuspended in fresh RPMI + 10% serum to a density of 500,000 cells per ml. The screened. For one 96 well plate, approximately 10 million cells (for 10 plates, 100 exact number of cells required will depend on the number of supernatants being million cells) are required. 12

Transfer the cells to a triangular reservoir boat, in order to dispense the cells into a 96 well dish, using a 12 channel pipette. Using a 12 channel pipette, transfer 200 ul of cells into each well (therefore adding 100, 000 cells per well).

channel pipette. In addition, a dose of exogenous interferon gamma (0,1 1.0, 10 ng) After all the plates have been seeded, 50 ul of the supernatants are transferred directly from the 96 well plate containing the supernatants into each well using a 12 is added to wells H9, H10, and H11 to serve as additional positive controls for the assay. ន

The 96 well dishes containing Jurkat cells treated with supernatants are placed in containing the remaining treated cells are placed at 40C and serve as a soutce of material pipette. The opaque plates should be covered (using sellophene covers) and stored atan incubator for 48 hrs (note: this time is variable between 48-72 hrs). 35 ul samples from each well are then transferred to an opaque 96 well plate using a 12 channel 200C until SEAP assays are performed according to Example 17. The plates for repeating the assay on a specific well if desired. 23 9

known to activate Jurkat T cells. Over 30 fold induction is typically observed in the As a positive control, 100 Unit/ml interferon gamma can be used which is positive control wells.

# Example 14: High-Throughput Screening Assay Identifying Myeloid Activity

The following protocol is used to assess myeloid activity by identifying factors, such as growth factors and cytokines, that may proliferate or differentiate myeloid cells. Myeloid cell activity is assessed using the GAS/SEAP/Neo construct produced in Example 12. Thus, factors that increase SEAP activity indicate the ability to activate the Jaks-STATS signal transduction pathway. The myeloid cell used in this assay is U937, a pre-monocyte cell line, although TF-1, HL60, or KG1 can be used.

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To transiently transfect U937 cells with the GAS/SEAP/Neo construct produced in Example 12, a DEAE-Dextran method (Kharbanda et. al., 1994, Cell Growth & Differentiation, 5:259-265) is used. First, harvest 2x10e<sup>7</sup> U937 cells and wash with PBS. The U937 cells are usually grown in RPMI 1640 medium containing 10% heatinactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 mg/ml streptomycin.

15 Next, suspend the cells in 1 ml of 20 mM Tris-HCl (pH 7.4) buffer containing 0.5 mg/ml DEAE-Dextran, 8 ug GAS-SEAP2 plasmid DNA, 140 mM NaCl, 5 mM KCl, 375 uM Na<sub>2</sub>HPO<sub>4</sub>.7H<sub>2</sub>O, 1 mM MgCl<sub>2</sub>, and 675 uM CaCl<sub>2</sub>. Incubate at 37°C for 45 min.

Wash the cells with RPMI 1640 medium containing 10% FBS and then

resuspend in 10 ml complete medium and incubate at 37°C for 36 hr.

The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400

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The GAS-SEAP/U937 stable cells are obtained by growing the cells in 400 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 400 ug/ml G418 for couple of passages.

These cells are tested by harvesting 1x10° cells (this is enough for ten 96-well plates assay) and wash with PBS. Suspend the cells in 200 ml above described growth medium, with a final density of 5x10° cells/ml. Plate 200 ul cells per well in the 96-well plate (or 1x10° cells/well).

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Add 50 ul of the supernatant prepared by the protocol described in Example 11.

Incubate at 37°C for 48 to 72 hr. As a positive control, 100 Unit/ml interferon gamma can be used which is known to activate U937 cells. Over 30 fold induction is typically observed in the positive control wells. SEAP assay the supernatant according to the protocol described in Example 17.

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# Example 15; High-Throughput Screening Assay Identifying Neuronal Activity.

When cells undergo differentiation and proliferation, a group of genes are activated through many different signal transduction pathways. One of these genes, EGR1 (early growth response gene 1), is induced in various tissues and cell types upon activation. The promoter of EGR1 is responsible for such induction. Using the EGR1 promoter linked to reporter molecules, activation of cells can be assessed.

Particularly, the following protocol is used to assess neuronal activity in PC12 cell lines. PC12 cells (rat phenochromocytoma cells) are known to proliferate and/or differentiate by activation with a number of mitogens, such as TPA (tetradecanoyl phorbol acetate), NGF (nerve growth factor), and EGF (epidermal growth factor). The EGR1 gene expression is activated during this treatment. Thus, by stably transfecting PC12 cells with a construct containing an EGR promoter linked to SEAP reporter, activation of PC12 cells can be assessed.

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The EGR/SEAP reporter construct can be assembled by the following protocol.

The EGR-I promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene 6:867-871 (1991)) can be PCR amplified from human genomic DNA using the following primers:

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# 5' GCGCTCGAGGGATGACAGCGATAGAACCCCCGG -3' (SEQ ID NO:6) 5' GCGAAGCTTCGCGACTCCCCGGGATCCGCCTC-3' (SEQ ID NO:7)

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Using the GAS:SEAP/Neo vector produced in Example 12, EGR1 amplified product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector using restriction enzymes Xhol/HindIII, removing the GAS/SV40 stuffer. Restrict the EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1 promoter.

25

To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30 dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and allowed to air dry for 2 hr.

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PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker) containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heatinactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and 100 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is done every three to four days. Cells are removed from the plates by scraping and resuspended with pipetting up and down for more than 15 times.

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Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine protocol described in Example 11. EGR-SEAP/PC12 stable cells are obtained by growing the cells in 300 ug/ml G418. The G418-free medium is used for routine growth but every one to two months, the cells should be re-grown in 300 ug/ml G418 for couple of passages.

To assay for neuronal activity, a 10 cm plate with cells around 70 to 80% confluent is screened by removing the old medium. Wash the cells once with PBS (Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as  $5 \times 10^5$ 

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Add 200 ul of the cell suspension to each well of 96-well plate (equivalent to 1x10<sup>5</sup> cells/well). Add 50 ul supernatant produced by Example 11, 37°C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ul of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 17.

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# Example 16: High-Throughput Screening Assay for T-cell Activity

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NF-kB (Nuclear Factor kB) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by expression of certain viral gene products. As a transcription factor, NF-kB regulates the expression of genes involved in immune cell activation, control of apoptosis (NF-kB appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

In non-stimulated conditions, NF- kB is retained in the cytoplasm with I-kB (Inhibitor kB). However, upon stimulation, I-kB is phosphorylated and degraded, causing NF- kB to shuttle to the nucleus, thereby activating transcription of target

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genes. Target genes activated by NF- xB include IL-2, IL-6, GM-CSF, ICAM-1 and

class 1 MHC.

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Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF-kB promoter element are used to screen the supernatants produced in Example 11. Activators or inhibitors of NF-kB would be useful in treating diseases. For example, inhibitors of NF-kB could be used to treat those diseases related to the acute or chronic activation of NF-kB, such as rheumatoid arthritis.

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To construct a vector containing the NF-kB promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF-kB binding site (GGGGACTTTCCC) (SEQ ID NO:8), 18 bp of sequence complementary to the 5' end of the SV40 early promoter sequence, and is flanked with an XhoI site:

10 5':GCGGCCTCGAGGGACTTTCCCGGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTTTCCGGGACTCTCAATTAG:3' (SEQ ID NO:9)

The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:

S':GCGGCAAGCTTTTTGCAAAGCCTAGGC:3' (SEQ ID NO:4)

the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2. (Stratagene) Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

25 3' (SEQ ID NO:10)

Next, replace the SV40 minimal promoter element present in the pSEAP2-promoter plasmid (Clontech) with this NF-xB/SV40 fragment using XhoI and HindIII. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

8

In order to generate stable mammalian cell lines, the NF-κB/SV40/SEAP cassette is removed from the above NF-κB/SEAP vector using restriction enzymes Sall and Notl, and inserted into a vector containing neomycin resistance. Particularly, the

gene, after restricting pGFP-1 with Sall and Notl. NF-xB/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP

the method for assaying supernatants with these stable Jurkat T-cells is also described wells H9, H10, and H11, with a 5-10 fold activation typically observed. in Example 13. As a positive control, exogenous TNF alpha (0.1,1, 10 ng) is added to created and maintained according to the protocol described in Example 13. Similarly, Once NF-kB/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are

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## Example 17: Assay for SEAP Activity

5 activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the Assay, and Reaction Buffers used below. following general procedure. The Tropix Phospho-light Kit supplies the Dilution, As a reporter molecule for the assays described in Examples 13-16, SEAP

dilution buffer into Optiplates containing 35 µl of a supernatant. Seal the plates with a heating. plastic sealer and incubate at 65°C for 30 min. Separate the Optiplates to avoid uneven Prime a dispenser with the 2.5x Dilution Buffer and dispense 15 µl of 2.5x

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prime with the Assay Buffer. Add 50 µl Assay Buffer and incubate at room Cool the samples to room temperature for 15 minutes. Empty the dispenser and

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time and start the second set 10 minutes later. takes about 10 minutes to read 5 plates on luminometer, one should treat 5 plates at each minutes. Since the intensity of the chemiluminescent signal is time dependent, and it table below). Add 50 µl Reaction Buffer and incubate at room temperature for 20 temperature 5 min. Empty the dispenser and prime with the Reaction Buffer (see the

25 results. An increase in chemiluminescence indicates reporter activity. Read the relative light unit in the luminometer. Set H12 as blank, and print the

### Reaction Buffer Formulation:

14	13	12	Ξ	10	# of plates
80	75	70	65	60	Rxn buffer diluent (ml)
4	3.75	3.5	3.25	<b>ω</b>	CSPD (ml)

50	49	48	47	46	45	44	43	42	4	40	39	38	37	36	35	34	33	32	31	30	29	28	27	26	25		23				19	18	. 17	16	15
260	255	250	245	240	235	230	225	220	215	210	205	200	195	190	185	180	175	170	165	160	155	150	145	140	135	130	125	120	115	. 110	105	100	95	90	85
13	12.75	12.5	12.25	12	11.75	11.5							9.75						8.25			7.5	7.25	7	6.75	6.5	6.25	6	5.75	5.5	5.25	5	4.75	4.5	4.25

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## Example 18: High-Throughput Screening Assay Identifying Changes in Small Molecule Concentration and Membrane Permeability

potential. These alterations can be measured in an assay to identify supernatants which sodium, pH, membrane potential, or any other small molecule which is detectable by a assay for calcium, this protocol can easily be modified to detect changes in potassium, Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane bind to receptors of a particular cell. Although the following protocol describes an fluorescent probe S

molecules. Clearly, any fluorescent molecule detecting a small molecule can be used The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small instead of the calcium fluorescent molecule, fluo-3, used here.

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96-well plate with clear bottom. The plate is incubated in a CO<sub>2</sub> incubator for 20 hours. For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash. 2

incubated at 37°C in a CO, incubator for 60 min. The plate is washed four times in the load the cells with fluo-3, 50 ul of 12 ug/ml fluo-3 is added to each well. The plate is A stock solution of 1 mg/ml fluo-3 is made in 10% pluronic acid DMSO. To Biotek washer with HBSS leaving 100 ul of buffer. 20

The tube is then placed in a 37°C water bath for 30-60 min. The cells are washed twice ul/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley CellWash with 200 ul, followed by an aspiration step to 100 ul final volume. re-suspended to 2-5x106 cells/ml with HBSS in a 50-ml conical tube. 4 ul of 1 mg/ml For non-adherent cells, the cells are spun down from culture media. Cells are fluo-3 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. with HBSS, resuspended to 1x10° cells/ml, and dispensed into a microplate, 100

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second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-3. The supernatant is added to the well, and a change in fluorescence is detected. To measure the fluorescence of intracellular calcium, the FLIPR is set for the following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 39

(6) Sample addition is 50 ul. Increased emission at 530 nm indicates an extracellular

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signaling even which has resulted in an increase in the intracellular Ca++ concentration

### Example 19: High-Throughput Screening Assay Identifying | Tyrosine Kinase Activity 'n

transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase ncluding the PDGF, FGF, EGF, NGF, HGF and Insulin receptor substamilies. In RPTK) group are receptors for a range of mitogenic and metabolic growth factors addition there are a large family of RPTKs for which the corresponding ligand is unknown. Ligands for RPTKs include mainly secreted small proteins, but also The Protein Tyrosine Kinases (PTK) represent a diverse group of membrane-bound and extracellular matrix proteins.

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receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members of which mediate signal transduction triggered by the cytokine superfamily of receptors Activation of RPTK by ligands involves ligand-mediated receptor dimerization, associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and noncytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor esulting in transphosphorylation of the receptor subunits and activation of the

2

kinase activity, the identification of novel human secreted proteins capable of activating yrosine kinase signal transduction pathways are of interest. Therefore, the following Because of the wide range of known factors capable of stimulating tyrosine protocol is designed to identify those novel human secreted proteins capable of ctivating the tyrosine kinase signal transduction pathways. 2

e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 00% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyne Silent Screen Plates purchased from 25

0% Matrigel purchased from Becton Dickinson (Bedford,MA), or calf serum, rinsed CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford,MA) are (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or with PBS and stored at 40C. Cell growth on these plates is assayed by seeding 5,000 alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, cells/well in growth medium and indirect quantitation of cell number through use of 3 35

plates can also be used in some proliferation experiments. used to cover the Loprodyne Silent Screen Plates. Falcon Microtest III cell culture

Loprodyne plates (20,000/200ml/well) and cultured overnight in complete medium To prepare extracts, A431 cells are seeded onto the nylon membranes of

- Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 and a cocktail of protease inhibitors (# 1836170) obtained from Boeheringer Mannheim 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na3VO4, 2 mM Na4P2O7 11, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example
- 5 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract (Indianapolis, IN) is added to each well and the plate is shaken on a rotating shaker for manifold and immediately placed on ice. To obtain extracts clarified by centrifugation Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum filtered through the 0.45 mm membrane bottoms of each well using house vacuum.
- 5 the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4°C at 16,000 x g.

methods of detecting tyrosine kinase activity are known, one method is described here Generally, the tyrosine kinase activity of a supernatant is evaluated by Test the filtered extracts for levels of tyrosine kinase activity. Although many

- 20 determining its ability to phosphorylate a tyrosine residue on a specific substrate (a a range of tyrosine kinases and are available from Boehringer Mannheim. PSK2 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and biotinylated peptide). Biotinylated peptides that can be used for this purpose include
- 25 components gently and preincubate the reaction mix at 30°C for 2 min. Initial the 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl2, 5 mM MnCl2, ATP/50mM MgCl2), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride. order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg2+ (5mM The tyrosine kinase reaction is set up by adding the following components in
- reaction by adding 10ul of the control enzyme or the filtered supernatant

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The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mm

mixture to a microtiter plate (MTP) module and incubating at 37°C for 20 min. This Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction

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POD(0.5w/ml)) to each well and incubate at 37°C for one hour. Wash the well as phospotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of antiallows the streptavadin coated 96 well plate to associate with the biotinylated peptide

tyrosine kinase activity. peroxidase activity is quantitated using an ELISA reader and reflects the level of absorbance of the sample at 405 nm by using ELISA reader. The level of bound incubate at room temperature for at least 5 mins (up to 30 min). Measure the Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and

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### Phosphorylation Activity Example 20: High-Throughput Screening Assay Identifying

20 5 Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other kinase activity described in Example 19, an assay which detects activation substituting these molecules for Erk-1 or Erk-2 in the following assay. phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase, phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other used. For example, as described below one particular assay can detect tyrosine (phosphorylation) of major intracellular signal transduction intermediates can also be As a potential alternative and/or compliment to the assay of protein tyrosine

25 3 rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4°C step can easily be modified by substituting a monoclonal antibody detecting any of the and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this are then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1 plate with 0.1ml of protein G (1ug/ml) for 2 hr at room temp, (RT). The plates are ther Specifically, assay plates are made by coating the wells of a 96-well ELISA

cultured overnight in growth medium. The cells are then starved for 48 hr in basal obtained in Example 11 for 5-20 minutes. The cells are then solubilized and extracts medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and

႘ filtered directly into the assay plate.

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After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody positive control, a commercial preparation of MAP kinase (10ng/well) is used in place (lug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac bound polyclonal antibody is then quantitated by successive incubations with background indicates a phosphorylation.

### Example 21: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

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these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in phenotype of interest (such as a disease) is be isolated. cDNA is then generated from RNA isolated from entire families or individual patients presenting with a seconds; 60-120 seconds at 52-58°C; and 60-120 seconds at 70°C, using buffer SEQ ID NO:X. Suggested PCR conditions consist of 35 cycles at 95°C for 30 solutions described in Sidransky, D., et al., Science 252:706 (1991).

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products analyzed to confirm the results. PCR products harboring suspected mutations polynucleotide kinase, employing SequiTherm Polymerase. (Epicentre Technologies). PCR products is then sequenced using primers labeled at their 5' end with T4 The intron-exon borders of selected exons is also determined and genomic PCR is then cloned and sequenced to validate the results of the direct sequencing. 8

PCR products is cloned into T-tailed vectors as described in Holton, T.A. and Graham, M.W., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals is identified by mutations not present in unaffected individuals. 25

Cg. et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the triphosphate (Boehringer Manheim), and FISH performed as described in Johnson, alterations in a gene corresponding to a polynucleotide. Genomic clones isolated Genomic rearrangements are also observed as a method of determining according to Example 2 are nick-translated with digoxigenindeoxy-uridine 5'corresponding genomic locus. 8 35

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propidium iodide, producing a combination of C- and R-bands. Aligned images for precise mapping are obtained using a triple-band filter set (Chroma Technology, Chromosomes are counterstained with 4,6-diamino-2-phenylidole and Brattleboro, VT) in combination with a cooled charge-coupled device camera

- (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson, Cv. Program System. (Inovision Corporation, Durham, NC.) Chromosome, alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and chromosomal fractional length measurements are performed using the ISee Graphical et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and S
  - translocations. These alterations are used as a diagnostic marker for an associated 으

## Example 22: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

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- a marker for a particular phenotype. Methods of detection are numerous, and thus, it is and if an increased or decreased level of the polypeptide is detected, this polypeptide is A polypeptide of the present invention can be detected in a biological sample, understood that one skilled in the art can modify the following assay to fit their particular needs.
- polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are described in Example 10. The wells are blocked so that non-specific binding of the coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method For example, antibody-sandwich ELISAs are used to detect soluble polypeptide to the well is reduced. 2 22

validate results. The plates are then washed three times with deionized or distilled water containing the polypeptide. Preferably, scrial dilutions of the sample should be used to The coated wells are then incubated for > 2 hours at RT with a sample to remove unbounded polypeptide.

concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at unbounded conjugate. 8

curve, using serial dilutions of a control sample, and plot polypeptide concentration on temperature. Measure the reaction by a microtiter plate reader. Prepare a standard Add 75 ul of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room 35

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Interpolate the concentration of the polypeptide in the sample using the standard curve. the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale).

## Example 23: Formulating a Polypeptide

S purposes herein is thus determined by such considerations. administration, and other factors known to practitioners. The "effective amount" for consistent with good medical practice, taking into account the clinical condition of the alone), the site of delivery, the method of administration, the scheduling of individual patient (especially the side effects of treatment with the secreted polypeptide The secreted polypeptide composition will be formulated and dosed in a fashion

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most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If and the interval following treatment for responses to occur appears to vary depending given continuously, the secreted polypeptide is typically administered at a dose rate of to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and polypeptide administered parenterally per dose will be in the range of about 1 µg/kg/day bag solution may also be employed. The length of treatment needed to observe changes continuous subcutaneous infusions, for example, using a mini-pump. An intravenous about 1 µg/kg/hour to about 50 µg/kg/hour, either by 1-4 injections per day or by to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject As a general proposition, the total pharmaceutically effective amount of secreted

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of administration which include intravenous, intramuscular, intraperitoneal, intrasternal patch), bucally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers subcutaneous and intraarticular injection and infusion. formulation auxiliary of any type. The term "parenteral" as used herein refers to modes to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or intraperitoneally, topically (as by powders, ointments, gels, drops or transderma administered orally, rectally, parenterally, intracistemally, intravaginally Pharmaceutical compositions containing the secreted protein of the invention are

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systems. Suitable examples of sustained-release compositions include semi-permeable al., J. Biomed. Mater. Res. 15:167-277 (1981), and R. Langer, Chem. Tech. 12:98-Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl methacrylate) (R. Lunger et copolymers of L-glutamic acid and gamma-ethyl-L-glutamate (Sidman, U. et al., Sustained-release matrices include polylactides (U.S. Pat. No. 3,773,919, EP 58,481), polymer matrices in the form of shaped articles, e.g., films, or mirocapsules. 105 (1982)), ethylene vinyl acetate (R. Langer et al.) or poly-D- (-)-3-hydroxybutyric The secreted polypeptide is also suitably administered by sustained-release

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EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese Pat. Appl. 83-118008; known per se: DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. USA 82:3688-369; polypeptides. Liposomes containing the secreted polypeptide are prepared by methods (1985); Hwang et al., Proc. Natl. Acad. Sci. USA 77:4030-4034 (1980); EP 52,322: acid (EP 133,988). Sustained-release compositions also include liposomally entrapped

is greater than about 30 mol. percent cholesterol, the selected proportion being adjusted for the optimal secreted polypeptide therapy. are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily, the liposomes

5 5 known to be deleterious to polypeptides. formulation preferably does not include oxidizing agents and other compounds that are employed and is compatible with other ingredients of the formulation. For example, the carrier, i.e., one that is non-toxic to recipients at the dosages and concentrations injectable form (solution, suspension, or emulsion), with a pharmaceutically acceptable formulated generally by mixing it at the desired degree of purity, in a unit dosage For parenteral administration, in one embodiment, the secreted polypeptide is

20 carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl of the recipient. Examples of such carrier vehicles include water, saline, Ringer's uniformly and intimately with liquid carriers or finely divided solid carriers or both. oleate are also useful herein, as well as liposomes. Then, if necessary, the product is shaped into the desired formulation. Preferably the Generally, the formulations are prepared by contacting the polypeptide

မ 23 ઝ enhance isotonicity and chemical stability. Such materials are non-toxic to recipients at poloxamers, or PEG immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or arginine; monosaccharides, citrate, succinate, acetic acid, and other organic acids or their salts; antioxidants such as the dosages and concentrations employed, and include buffers such as phosphate, polyarginine or tripeptides; proteins, such as serum albumin, gelatin, or ascorbic acid; low molecular weight (less than about ten residues) polypeptides, e.g., sorbitol; counterions such as sodium; and/or nonionic surfactants such as polysorbates. manose, or dextrins; chelating agents such as EDTA; sugar alcohols such as mannitol or disaccharides, and other carbohydrates including cellulose or its derivatives, glucose, The carrier suitably contains minor amounts of additives such as substances that

concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of The secreted polypeptide is typically formulated in such vehicles at a

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about 3 to 8. It will be understood that the use of certain of the foregoing excipients, carriers, or stabilizers will result in the formation of polypeptide salts.

into a container having a sterile access port, for example, an intravenous solution bag or Sterility is readily accomplished by filtration through sterile filtration membranes (e.g., 0.2 micron membranes). Therapeutic polypeptide compositions generally are placed Any polypeptide to be used for therapeutic administration can be sterile. vial having a stopper pierceable by a hypodermic injection needle.

formulation for reconstitution. As an example of a lyophilized formulation, 10-ml vials resulting mixture is lyophilized. The infusion solution is prepared by reconstituting the are filled with 5 ml of sterile-filtered 1% (w/v) aqueous polypeptide solution, and the Polypeptides ordinarily will be stored in unit or multi-dose containers, for example, sealed ampoules or vials, as an aqueous solution or as a lyophilized lyophilized polypeptide using bacteriostatic Water-for-Injection.

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pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of present invention may be employed in conjunction with other therapeutic compounds. The invention also provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical

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# Example 24: Method of Treating Decreased Levels of the Polypeptide

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administering the polypeptide of the present invention, preferably in the secreted form. Thus, the invention also provides a method of treatment of an individual in need of an pharmaceutical composition comprising an amount of the polypeptide to increase the It will be appreciated that conditions caused by a decrease in the standard or increased level of the polypeptide comprising administering to such an individual a normal expression level of a secreted protein in an individual can be treated by activity level of the polypeptide in such an individual.

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polypeptide is in the secreted form. The exact details of the dosing scheme, based on For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the polypeptide for six consecutive days. Preferably, the administration and formulation, are provided in Example 23.

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# Example 25: Method of Treating Increased Levels of the Polypeptide

a polypeptide, preferably a secreted form, due to a variety of etiologies, such as cancer. present invention. This technology is one example of a method of decreasing levels of Antisense technology is used to inhibit production of a polypeptide of the

2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, For example, a patient diagnosed with abnormally increased levels of a provided in Example 23.

## Example 26: Method of Treatment Using Gene Therapy

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hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to separated into small pieces. Small chunks of the tissue are placed on a wet surface of a expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and issue culture flask, approximately ten pieces are placed in each flask. The flask is urned upside down, closed tight and left at room temperature over night. After 24 One method of gene therapy transplants fibroblasts, which are capable of he bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS,

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penicillin and streptomycin, is added. The flasks are then incubated at  $37^{\circ}\mathrm{C}$  for approximately one week. 2

At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is erminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and fractionated on agarose gel and purified, using glass beads. 25

using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear The cDNA encoding a polypeptide of the present invention can be amplified backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to 3 33

the purpose of confirming that the vector has the gene of interest properly inserted transform bacteria HB101, which are then plated onto agar containing kanamycin for

calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is packaging cells are now referred to as producer cells). packaging cells now produce infectious viral particles containing the gene (the then added to the media and the packaging cells transduced with the vector. The culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% The amphotropic pA317 or GP+am12 packaging cells are grown in tissue

2 5 titer of virus is high, then virtually all fibroblasts will be infected and no selection is containing the infectious viral particles, is filtered through a millipore filter to remove media is harvested from a 10 cm plate of confluent producer cells. The spent media, infected, the fibroblasts are analyzed to determine whether protein is being produced. selectable marker, such as neo or his. Once the fibroblasts have been efficiently required. If the titer is very low, then it is necessary to use a retroviral vector that has a removed from a sub-confluent plate of fibroblasts and quickly replaced with the media detached producer cells and this media is then used to infect fibroblast cells. Media is from the producer cells. This media is removed and replaced with fresh media. If the Fresh media is added to the transduced producer cells, and subsequently, the

after having been grown to confluence on cytodex 3 microcarrier beads. The engineered fibroblasts are then transplanted onto the host, either alone or

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therefore, are within the scope of the appended claims variations of the present invention are possible in light of the above teachings and described in the foregoing description and examples. Numerous modifications and It will be clear that the invention may be practiced otherwise than as particularly

hereby incorporated herein by reference disclosures) in the Background of the Invention, Detailed Description, and Examples is applications, journal articles, abstracts, laboratory manuals, books, or other The entire disclosure of each document cited (including patents, paten

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3 8 35 ဗ 23 20 15 5 S (1) GENERAL INFORMATION: (iii) NUMBER OF SEQUENCES: 644 (ii) TITLE OF INVENTION: 186 Human Secreted Proteins (i) APPLICANT: (v) COMPUTER READABLE FORM: (iv) CORRESPONDENCE ADDRESS: (vi) CURRENT APPLICATION DATA: ਉ (C) CITY: Rockville (D) SOFTWARE: ASCII Text (C) OPERATING SYSTEM: MSDOS version 6.2 (B) COMPUTER: HP Vectra 486/33 (A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage  $\Xi$ (B) STREET: 9410 Key West Avenue (A) ADDRESSEE: Human Genome Sciences, Inc. (A) APPLICATION NUMBER (F) ZIP: 20850 (C) CLASSIFICATION: (B) FILING DATE: March 6, 1998 COUNTRY: USA STATE: Maryland Human Genome Sciences, Inc. et al

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(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER:

(B) FILING DATE:

10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Eq.   10   MARIE A. Andreas Brookes, Ed.   10   MARI	-			
10 SEQUENCY CHARGEST STATES   10	(viii) ATTORNEY/AGENT INFORMATION:		(2) INFORMATION FOR SBQ ID NO: 2:	·· · · · · · · · · · · · · · · · · · ·
(ii) SECURITY SECTION 10: 10: 10: 10: 10: 10: 10: 10: 10: 10:	(A) NAME: A. Anders Brookes, Esq.	8	ğ	
10   Tap Set Yea Tap Set	(B) REGISTRATION NUMBER: 36,373		(B) TYPE: amino acid (D) TOPOLOGY: linear	
15   17   17   18   18   18   18   18   18	(C) REFERENCE/DOCKET NUMBER: PS002.PCT		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:	
15   (2) INSTRUMENTION FOR SEQ ID NO; 3;   (1) SEQUENCE CHANATTEGETICS;   (1) A LENGTH is the base pairs   (1) A LENGTH is the base pairs   (2) STRUMENTISS; duable   (2) STRUMENTISS; duable   (2) STRUMENTISS; duable   (2) STRUMENTISS; duable   (3) TOPUCLOT; Linear   (4) SEQUENCE DISSORIFTION; SEQ ID NO; 3;   (2) A LENGTH TOCCOLANT CHATCOCO ANTOLOTY CHATCOCO ANTOLOTY CHATCOCO ANTOLOTY CHATCOCO ANTOLOTY CHATCOCO ANTOLOTY CHATCOCO ANTOLOTY; LINEAR   (3) TOPUCLOTY; LINEAR   (4) CONTRACTOCOLATY; LINEAR   (5) TOPUCLOTY; LINEAR   (6) TOPUCLOTY; LINEAR   (7) TOPUCLOTY; LINEAR   (8) TOPUCLOTY; LINEAR   (9) TOPUCLOTY; LINEAR   (1) SEQUENCE CHANACTERISTICS;   (1) SEQUENCE CHANACTERISTICS;   (1) SEQUENCE CHANACTERISTICS;   (1) SEQUENCE CHANACTERISTICS;   (2) STRUMENTERISTICS;   (3) TOPUCLOTY; LINEAR   (4) TOPUCLOTY; LINEAR   (5) TOPUCLOTY; LINEAR   (6) TOPUCLOTY; LINEAR   (7) TOPUCLOTY; LINE			Š	
(i) INFORMATION FOR SEQ ID NO; 3;  (ii) SEQUENCE CHANACTERISTICS;  (iii) TIPE: INCLUDENCE CHANACTERISTICS;  (iv) TIPE: INCLUDENCE CHANACTERISTICS;  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TERRIBATICS; double  (iv) TERRIBATICS; double  (iv) TERRIBATICS; double  (iv) TERRIBATICS; double  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) SEQUENCE DESCRIPTION; SEQ ID NO; 4;  (iv) SEQUENCE DESCRIPTION; SEQ ID NO; 4;  (iv) SEQUENCE DESCRIPTION; SEQ ID NO; 5;  (iv) SEQUENCE CHANACTERISTICS;  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) TOPICLOCY; Linear  (iv) SEQUENCE DESCRIPTION; SEQ ID NO; 5;	(vi) TELECOMMUNICATION INFORMATION:			
(1) INPORMATION FOR SEQ ID NO; 3;  (1) SEQUENCE CHARACTERISTICS;  (1) TIPES INCLISE CHARLE  (1) TIPES INCLISE CHARLE  (2) TIPES INCLISE CHARLE  (3) COCCOUNTING ATTOCCCY; Linear  240  (4) COCCOUNTING TOWN SEQ ID NO; 4;  (4) SEQUENCE DESCRIPTION; SEQ ID NO; 4;  (5) INPORMATION FOR SEQ ID NO; 4;  (6) TIPES INCLISE CHARLE  (7) TIPES INCLISE CHARLE  (8) (1) SEQUENCE CHARACTERISTICS; (9) LEAVE CHARLES	(A) TELEPHONE: (301) 309-8504	115		
(i) SEQUENCE CHARACTERISTICS:  (ii) LABOTHIN (B. Due putre  (ic) STRANDENDESS: double  (ic) STRANDENDESS: double  (ic) STRANDENDESS: double  (ic) STRANDENDESS: double  (ic) STRANDENDESS: double  (id) SEQUENCE DESCRIPTION: SEQ ID NO: 3:  (id) SEQUENCE DESCRIPTION: SEQ ID NO: 4:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) TOPICLOST: Jinear  (id) TOPICLOST: Jinear  (id) TOPICLOST: Jinear  (id) TOPICLOST: Jinear  (id) TOPICLOST: Jinear  (id) TOPICLOST: Jinear  (id) TOPICLOST: Jinear  (id) SEQUENCE DESCRIPTION: SEQ ID NO: 4:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE CHARACTERISTICS:  (id) SEQUENCE DESCRIPTION: SEQ ID NO: 5	(B) TELEFAX: (301) 309-8439		(2) INFORMATION FOR SEQ ID NO: 3:	
25   (x1) SEQUENCE DESCRIPTION; SEQ ID NO; 3;	(2) INPORMATION FOR SEQ ID NO: 1:	20	SEQUENC (A) (B) (C) (D)	
CCCGAMATHY CTGCCCGAMATHY TCCCCGAMAT GATTTCCCCC AMPGATTTCC   120	(i) SEQUENCE CHARACTERISTICS: (a) IDATH: 711 base pairs	25	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 3:	
120   35   (2) INPORMATION FOR SEQ ID NO: 4:     120   35   (2) INPORMATION FOR SEQ ID NO: 4:     120   35   (1) SEQUENCE CHARACTERISTICS:     120   400   (1) SEQUENCE CHARACTERISTICS:     120   400   (2) TRANSENSES: double     120   420   (2) INPORMATION FOR SEQ ID NO: 4:     120   420   (2) INPORMATION FOR SEQ ID NO: 5:     120   600   (1) SEQUENCE CHARACTERISTICS:     120   (2) INPORMATION FOR SEQ ID NO: 5:     121   (2) INPORMATION FOR SEQ ID NO: 5:     122   (3) INPORMATION FOR SEQ ID NO: 5:     123   (4) INPORMATION FOR SEQ ID NO: 5:     124   (1) SEQUENCE CHARACTERISTICS:     125   (2) INPORMATION FOR SEQ ID NO: 5:     125   (3) INPORMATION FOR SEQ ID NO: 5:     126   (4) INPORMATION SEQ ID NO: 5:     127   (4) INPORMATION SEQ ID NO: 5:     128   (4) INPORMATION SEQ ID NO: 5:     129   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     120   (4) INPORMATION SEQ ID NO: 5:     121   (5) INPORMATION SEQ ID NO: 5:     121   (6) INPORMATION SEQ ID NO: 5:     121   (6) INPORMATION SEQ ID NO: 5:     121   (6) INPORMATION SEQ ID NO: 5:     121   (6) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     121   (7) INPORMATION SEQ ID NO: 5:     122   (7) INPORMATION SEQ ID NO: 5:     123	(B) TYPE: mucleic acid (C) STRANDENESS: double	·	COCCTCGAG ATTICCCCGA AATCTAGATT TCCCCGAAAT GATTTCCCCG AAATGATTTC	09
120   35   13FORMATION FOR SEQ ID NO: 4:     120   35   (1) INFORMATION FOR SEQ ID NO: 4:     120   35   (1) SEQUENCE CHARACTERISTICS:     120   (1) LENGTH: 27 base pairs     120   (2) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) LENGTH: 27) base pairs     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) LENGTH: 27) base pairs     120   (5) TOPOLOGY: 1 innear     120   (6) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (8) TOPOLOGY: 1 innear     120   (9) TOPOLOGY: 1 innear     120   (1) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) TOPOLOGY: 1 innear     120   (5) TOPOLOGY: 1 innear     120   (6) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (8) TOPOLOGY: 1 innear     120   (9) TOPOLOGY: 1 innear     120   (1) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) TOPOLOGY: 1 innear     120   (5) TOPOLOGY: 1 innear     120   (6) TOPOLOGY: 1 innear     120   (6) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (8) TOPOLOGY: 1 innear     120   (9) TOPOLOGY: 1 innear     120   (1) TOPOLOGY: 1 innear     120   (2) TOPOLOGY: 1 innear     120   (3) TOPOLOGY: 1 innear     120   (4) TOPOLOGY: 1 innear     120   (5) TOPOLOGY: 1 innear     120   (6) TOPOLOGY: 1 innear     120   (6) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOPOLOGY: 1 innear     120   (7) TOP	(D) TOPOLOGY: linear	•	COCGNANTAL CIGCCATCIC AATTAG	98
120   35   (2) INFORMATION FOR SEQ ID NO: 4:	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:	30		
120   35	GGGATCGGA GCCCAAATCT TCTGACAAAA CTCACACATG CCCACGGTGC CCAGCACCTG	09		
180	CGAGGG TECACCOTCA GTCTTCCTCT TECOCECAAA ACCCAAGGAC ACCCTCATGA		(2) INFORMATION FOR SEQ ID NO: 4:	
100   170PE: nucleic acid   (1) TYPE: nucleic acid   (1) TYPE: nucleic acid   (1) TYPE: nucleic acid   (1) TYPE: nucleic acid   (2) TYPE: nucleic acid   (3) TYPE: nucleic acid   (4) LENGTH: 271 base pairs   (2) TYPE: nucleic acid   (3) TYPE: nucleic acid   (4) LENGTH: 271 base pairs   (5) TYPE: nucleic acid   (5) TYPE: nucleic acid   (5) TYPE: nucleic acid   (6) TYPE: nucleic acid   (7) TYPE	CCGGAC TCCTGAGGTC ACATGCCTGG TGGTGGACGT AAGCCACGAA GACCCTGAGG		SEO	
100   100 TOPOLOGY: 1 Linear   1,000   1,000 TOPOLOGY: 1 Linear   1,000   1,	GITCAA CTGGTACGTG GACGGCGTGG AGGTGCATAA TGCCAAGACA AAGCCGGGG		(B) TYPE: nucleic acid (C) STRANDEIMESS: double	
420 420 420 420 420 420 420 420 480 480 480 540 50 600 61) INFORMATION FOR SEQ ID NO: 5: 600 (1) SEQUENCE CHARACTERISTICS: 600 (1) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 5: (3) LENGTH: 271 base pairs (4) LENGTH: 271 base pairs (5) TRANDEDNESS: double (C) STRANDEDNESS: double (C) STRANDEDNESS: double (C) STRANDEDNESS: double (C) STRANDEDNESS: double (C) STRANDEDNESS: double (C) STRANDENESS: dou	SCASTA CAACAGCACG TACCGTGTGG TCAGCGTCCT CACCGTCCTG CACCAGGACT	٠		
420 45 460 540 600 600 (2) INFORMATION FOR SEQ ID NO: 5: 660 (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 271 base pairs (B) TYPE: nucleic acid (C) STRANDENNESS: double (C) S	GANTGG CANGGAGTAC AMOTGCAAGG TCTCCAACAA AOCCCTCCCA ACCCCCATCG	360	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 4:	
540 540 550 (1) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 5: (3) LEWATH: 271 base pairs (4) LEWATH: 271 base pairs (B) TYPE: INCISIC acid (C) STRANDEDNESS: double (B) TOPOLOGY: Linear (C) TOGAGATIT CCCCGAAATC TAGATITCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG	MACCHT CTCCAAAGCC AAAGGCAGC CCCGAGAACC ACAGGTGTAC ACCCTGCCCC		GCGCCAGCT TTTTGCAAAG CCTAGGC	27
600 (1) INFORMATION FOR SEQ ID NO: 5: 600 (i) SEQUENCE CHARACTERISTICS: 660 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 271 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) STRANDEDNESS: double (D) STRANDEDNES	COOGGGA TGAGCTGACC ANGAACCAGG TCAGCCTGAC CTGCCTGGTC ANAGGCTTCT	480		
660 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 271 base pairs (A) LENGTH: 271 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (C) TOPOLOGY: linear (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:  CTCCAGATTT CCCCGAAATC TAGATTTCCCCGAAAT GATTTCCCCG	завста ситоососто састоссава сслатоосса ососвавалс ластислава		(2) INFORMATION FOR SEQ ID NO: 5:	
SCAGCAG GGGAACGTCT TCTCATGCTC COTGATGCAT GAGGCTCTGC 660  (B) TYPE: nucleic acid (C) STRANDEDNESS: double SCAGAAG AGCCTCTCC TGTCTCCGGG TAAATGAGTG CGACGCCCC 720  (Ai) SEQUENCE DESCRIPTION: SEQ ID NO: 5:  (TCGAGATTT CCCGAAATC TAGATTTCCC CGAAATGATT TCCCGAAATC TAGATTTCCC CGAAATGATT TCCCGAAATC GATTTCCCCG	SOCTEC COTOCTOGAC TECCACGOCT CETTETTECT CTACAGEAAG CTCACCGTGG			
SCAGAAG AGCOTOTOCOGGG TAAATGAGTG CGAGGGCCCC 720 733 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:  CTCGAGAATT CCCCGAAATC TAGATTTCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG	NAGCAG GIGGCAGCAG GGGAACGTCT TCTCATGCTC CGTGATGCAT GAGGCTCTGC	099	E	
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:  CTCGAGAITT CCCCGAAAITC TAGAITTCCC CGAAAITGAIT TCCCCGAAAIT GAITTCCCCG	ICCACTA CAGGCAGAAG AGCCTCTCCC TGTCTCCGGG TAAATGAGTG CGAGGGCCGC			
CTCGAGATTT CCCCGAAATC TAGATTTCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG	CCTAGAG GAT	733	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 5:	
		09	CTCGAGATIT CCCCGAAATC TAGATITCCC CGAAATGATT TCCCCGAAAT GATTTCCCCG	09

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420	AGAAAAAAC TOOCITCATI TCTOTGAAAT TOCTCTTTGA AAATTTCTTT TTACACOTGT	60	(1) SEQUENCE CHARACTERISTICS:	60
360	GATATOSTAA TITTCATAAC AGATSICAST TITGAACCAA GAATTGSTGA TITGTTTATA			
300	ТОПАЛАГИДА СТГАСТТОСА СТГТСАЛСАТ САСТТАСТСС ТТОТСАТСТТ АСАЛАГИТСТ	,		ડ
240	TITIMATCIT TOTATATTIT TYMAMATCC TITICICCIC ATCATIOCCT TITITIOTOGI	12	GOGGACTITIC CC	
180	ANICACITYTY TOTOTYTIAN COTOTAACCA AAAAATIGIT TAATTYYOGA TOOCAAANGT		(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 8:	
120	GCAAGCTGTT AAAGATCTTG GATCCCATTA TATAGTATGT ATAGCTGAAA TCTGTAATTC	. 50		50
60	GGCACGAGGT AATTYCTACC AGAAATTYCC AGAGCATTAT GTAGGTAGAA AAAAATGCAA		(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 11:	<b>4</b> 5	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 12 base pairs	45
	(B) TYPE: nucleic acid (C) STRANDELNESS: double (D) TOPOLOGY: linear		(2) INFORMATION FOR SEQ ID NO: 8:	
	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 582 base pairs	40		40
	(2) INFORMATION FOR SEQ ID NO: 11:	31	GCGAAGCTTC GCGACTCCCC GGATCCGCCT C	
		35	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 7:	35
256	CTTTTOCAAA AAGCTT			
240	OCCOCCTOS COCTOTARIOS TATTOCADAA GIAGIGAAGGA COCTITITITIS GAGOCCIAGO	, c	(A) LENGTH: 31 base pairs (B) TYPE: nucleic acid	30
180	CAGTICCOCC CATTCICCOC CCCATOGCTG ACTAATITIT TITATITATIG CAGAGGCCGA	30	(i) SEQUENCE CHARACTERISTICS:	
120	CANTINGTEA GENECIATAG TECEGECECT ANCIECEGECE ATECEGECEC TANCIECEGEC		(2) INFORMATION FOR SEQ ID NO: 7:	
60	CYCHADOGGA CTTTCCCOOG GACTTTCCCG GGACTTTCCA TCTGCCATCT	. 25		25
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:	32	осостсанов антинсавов атнамассес ов	
	(C) STRANDERWESS: double (D) TOPOLOGY: linear	20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:	20
		3	(B) TYPE: nucleic acid (C) STRANDENNESS: double (D) TOPOLOGY: linear	
	(2) INFORMATION FOR SEQ ID NO: 10:	15		15
			(2) INFORMATION FOR SEQ ID NO: 6:	
73	CONTENANT TAG	10		10
60	geogetical occaverine ecosocient necesscien inecessien inechnecie	271	TITIOGAGGC CTAGGCTITT GCAAAAAGCT T	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:	240	TTATIOCAGAG GEOGAGGEEG CETEGGEETE TGAGETAFTE CAGAAGTAGT GAGGAGGETT	v
		180	GOODCETAACT COGCOCAGET COGCOCAFTC TOCGCOCCAT GGCTGACTAA TETETETEAT	
	(A) LENOTH: 73 base pairs (B) TYPE: nucleic acid	120	ANATHRICING CARCICARTE AGREAGEAGE CARRIGICEG COCCENACIO, OCCCEATOCO	
	244 ,		243	
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	. 245		246
	AAGCCAACTG AGATACCGTG ATGGTOTTGA TITTCTTTCAA TGATGCTTAC CATCTATTTT	480	ACCCTAGCAM TTGGGGGGTG AGCGGGTGGA TCACGAGGTC AGGAGATCGA GACCATCCTG
v	AGCCACTGAG CCTTTTATTA TTTGTCTATT TGTAAAGTTT ATTTGTCTTA ACTCATTTAA	540	gstarcatigg tgaaaccccg tctctactaa aaatacaaaa aaaaaaaaa aaaa
1	taaatatact gettaectgt tectgaaaaa aaaaaaaaa aa		
9		<u>s</u>	(2) INFORMATION FOR SEQ ID NO: 14:
2	(2) INFORMATION FOR SEQ ID NO: 12:		(i) SEQUENCE CHARACTERISTICS: (a) Instant: 314 hase mairs
	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 465 base pairs		(C) STRANDENESS: double
15	(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear		(D) TOPQLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:
ć	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:		TIATGITGGG GAGCAAGACC EGATAGCCAG CCTTTACATG GGAGTATAAT TCTGTCCTCC
3	OTTOGGGGG GAGCCGAGC TOCTOCGGGG CTTCGTCGCC GCCCAGACA CACCTACTCG	09	ATCTCATAAG CCCCAGTACC TGAGCCAGAA TGATTATAAC CAACCACAC GTCTCTTTAT
	CACGGCCGCC GCGCCTGGCT ATGATGTTCC TCACCCAGGG CGGCCTCTG CCCTCTACTC	120	CATGGATGGC TITAGCAGTA GGTTATTITC ATCATTGCCA ITTGTAGCTC TACAGTGGTT
25	отоссмостс систтоссмо всмоглассс тесесмост ттемогаетта стехнател	180 25	TATAGTAATT TCTCATCTTT TAAGTCTCTC CCTCAGTGCC TGTTGTTATC AAACTCATTG
	CCTOTTIGGA TIGACTAAAA GGACCTTIOT GTOGGAACAG GTGCTCCCCA AACACCTICC	240	CICTCTCANG CAGITGAGCT CTGCATTCTC CCTTATGGGG GAGAGCTGTG TTGGAGAGAG
90	TIGHTIGGETIGE CAGGEAGGE CALAGGGGCAG CACTEATEAG CACCTECETTG	300	AGAATATNAC TTCC
20	GACCCCTGCA GGGCAGGCAG CTTGGGGCCG AGCCCAAGGA TTTGGCTCTG CTGCCCCAA	360	
	GOGGACAGGA AGCOTOTTOG GCOTOTTOCC TTCCTOGACA AGGOCCOCTG COTTTGCOTC	420	(2) INPORMATION FOR SPO ID NO: 15:
35	ACATAAACTG TACAGTATTT TCATTAAAAG CCTCTTTCAT AAAAA	465 35	(1) PERMITANTA MINERAMODICANTO.
Ş		04	(1) SEQUENCE CHARACIEALS:  (A) LENGTH: 613 base pairs  (B) TTPE: nucleic acid  (C) STRANDEDRESS double
<del>5</del>	(2) INPORMATION FOR SEQ ID NO: 13:	F	(v) ICPOLOUS: ASSESSED TO NO. 15.
ų		¥.	CICATATTGC COTCIGGCTA AAAGTGAACA TGCCATTGAT CAATCTGCTT TTATTATATT
G	(C) STRANDENNESS: double (D) TOPOLOGY: linear	<del>,</del>	ATGITCCTAA TOOTGCCAAG CAAGACAAGA AGTAGAAGA AAGATGOTOT AAGCTCAAGA
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:		ACCCACTAAA TCTATCCTAT GGCCTGGGTT CACCCAGCCT GCTTTGTGGA TTTTGTCTCA
20	ATOCAATTCC TGCTCACAGC CTTTCTGTTG GTGCCACTTC TGGCTCTTTG TGATGTCCCC	09	CININACAGA GCICCCAAGG AGACTGCAGA GICAGCTCCC TTAAGCACTG TAACTAAAAGC
	ATAICCCTAG GCITCTCCC CTCCTAGAAG GGCITCTTGA TAGAITAGAA AATAAGAATG	120	CHARCTETIC COTTCCACC AACAATOTYC CCAGCTCATC CTCTTTCCCR AAGTCCCCTT
ť	AGTGACATIT CCTATGTGCA TATAAGAAGG AGCCACAAGA CATGTCTTTT AAATAAAAGG	180 55	TCTGCCCCAG ATGCGAATTG CATTTAACTA ATCCTCAAGT GAAATGTCCA CACAGRATTC
3	ACAGNOTOCOA TCCTTTTAGC TGCCGAATAG AACCTTGGTC TCATCCTCCT GGAGCTAGGC	240	CATTITIAATT AGCATACCAT AGTITITIGIG CAAATTIOCT TTCAGARGAC TCCCATTGCA
	CITTAAAACA GCITICIGIGI FICICATITIG ICTCAGIGIT TIGGCAGGGI FITAATCGGAA	300	GCTGCTCAGA GACGCTAAWG GCAGGGCCTC TTGAWGCTTT CCCGATAGCT TTCAGCTGCA
99	AGAIDAIGIT COSTITAAAA IATITCCIAA TGAGGCGGG GGTGGTGGCT CACGCCTGTA	980	ATROCTETTA GOCAGAATGE CATGAGOSTE CTGCCCAACT STATTACTOG GGAACACCTG

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(2) INFORMATION FOR SEQ ID NO: 18:

550	ааалалаа		
540	ATGRICCCIAT GGGCCCTGGA GGCRGCCCAT TRANGCATCT GGCTCGTTTT TGGRARARA		414
480	TOTOGRATIOC TORROTICARC CARRIECTEG CARRICTORG OCCICATITA AGGGRITICIG	, ,	360
420 .	AGGACCTTGC TACTOCOCAA GCCAGAGCCC ATCAGCCAGG CCTGCTGTGA GCCACCTGCC		300
360	астоотосью отвоествые товечается отестьюмые отетствамс атетовыств	50	240
300	GEOCRITIATE CEAGGAAACT TEATGETITE TAGAAGCEAA GEAGCIGCIG GEACTEAGGE		180
240	AGAGECCEAG CETOTOCTGA ATTGACATCA GTGETTCCCT GAACTGCCTC CCCCACCCCT		120
180	GRECHOSONY OCHOCOTOGA OCTOOCOTOC TYGITICTICOG CCTOCTOCTG CCCCCACCCC	45	60
120	тистосносе товносткое тоннованиее тоснетскогт остоссиите скозстисто		
60	CCCCCCCCC CCCCACACT TICHGGAGTC ACCCCCAGC ATTIGGGGTT GGGTTGGCCC	40	
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 19:		
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 550 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: double  (D) TOPOLOGY: linear	35	
	(2) INFORMATION FOR SEQ ID NO: 19:	30	356
			300
469	ACTIVITOTOT ATTOTTOCTC ATACTITOCCT ACCCCCAAAT TAATATCAG	25	240
420	GATOTICOTOT TOGAGGGATG AAGATOGAGT TATICCTICG AGAAATTICCT AGACGCCTTC		180
360	GATTTACCTT AAGGTCCAAA CCTCAGAACC CTCGGGCACC TGAGAGAGAT GTTTTGCGGC	•	120
300	оссьювансе аппанамнося ссмымансет поменанана оснанисска андиалисьс	20	60
240	отпоасста стгоссосов выгомсост астгеттесы вывыссымая оботсосттт		
180	CTTCATHOTC TICAGGCTGA GGGAGGTCTG GCAAGCCTTG CCCCTCATTT TGTTTGCGGT	15	
120	садваасуте дамогдатов тетоттесте естототоме атадотовая тактелеесе		
. 60	AATCACCATT GCAATACAAA TGATCTGCCT GGTGAATGYT GAGCTGTACC CCACATTCGT		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:		
	(1) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 469 base pairs  (B) TYPE: nucleic acid  (C) STRANDEINNESS: double  (D) TOPOLOGY: linear	5	613
	And designation with the Man and the same an		600

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GAAACTANAT CCCGGGGCTT TTAACNOGTA CTTGGGAAAT AAGTATTGGG TAATCACTAA GNGGACATTG ACTGCACCAA ACCAAAGCTA TAGAAAGAAA TGATTGACTT TTTAAAATAT

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:

8

ATTCACATTA ACTOTOCTAG GATACTICIC TIGAGGCITI GGAAAACTIC TICCTIGAAA
TITIGCATATC CACTICGAGTI CIGICACCAA AGATITITAAT CTICAGATOG CAATTITCCIC

TCTCCCAGAA AAAAGTACTA CAACAGGCTC AAGGGATATG CTTTOGTGGT CAAGGGATTA

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CACTATIGGTT TECCTTETGT TEACAATGGT ATTTACAGGA GACCTTGTCA TEAGAGGAGG
TACTGAACTA TETTTATGAC TTTGGATTTG ATCAGAGGTT TAAAAAAAAA AAAA

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CCCTGAAAAC GTAATTTGTG GTGTTACCAA AGGACCACAG GGGAAAAAAA AAAAAA

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(2) INFORMATION FOR SEQ ID NO: 17:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 414 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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OTGAOTOGIC TOCCTOCCA AGGAOCCIGA TIGGIGGGAA AUGGCATCAT CIAATAIGAT

TIMITITIMO TIGOAGAIGA COAGGICICO COCCOACAGO CICIGIOTOGO ICCOTOATIG

GOGAAGCCAT TIGGICCIGG TIAIGITTAT TACAACATCA TIGCACICIG GOACICCAGI

ссссссски талассства оспоталала ттитисств тапазатся тепатапасс осстоята такисска спестатос лалагасае слосслатся таласске

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

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(2) INFORMATION FOR SEQ ID NO: 16:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 356 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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CCAMATATICC CAC

ATTOGCTAGA AGITGATICET CCTGTAACTT TICTGAGTIC TITACATITA CTCGTGAAAC

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			ACTCAGGTTG AAGGAAATAT ACATAAATAA GATAAAGCTG ACCTGTAGAT ATAGCAGGTT
	(2) INFORMATION FOR SEQ ID NO: 20:		ATAMAGCITA GAOTTOICIA AGTIGAOTOC AMAITITICCI CIGATCITIC IGAIGCOGA
2	(i) SEQUENCE CHARACTERISTICS:	\$	
	(A) LENGTH: 741 base pairs		CAAAAAAGCA GTCATGTTTG TTATGTGATT GGAATGGAAC CCGAGAAGAG AGCATGCTGT
	(B) TYPE: nucleic acid (C) STRANDEMPSS: double		GITCTIGIGG GACAGGAAAG CITGCGIGCA CCAAGICIGA ACCACCACCI TCATGGTGAC
9			AND AND THE CONTRACT STATEMENT AND STATEMENT STREET, S
2	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 20:		GCAGTGGAAA CGSTCATCTTT TTTTTCCCCTCA CAGTGGTATTTT TTTTTCCAAAAA CGSTCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA
	TCTTGAAGAG TGTACAGTAC AGGATTATTA TAATGAAAGT TTATATCAAC AGGGTTTCGT	09	TOCTOCTABA ACACACTIC CANCETAANG PARCETOCOTA
15	TOCCTCTGCA TATATTATAA GCAAAAGAGA TTGGTAAAGT GCCACAGTAT TCCAGATAAC	120 150	AACAGCCTTT GOGAGGTCCT CCTTGATTCA TSGATGAAAC CTGGAACATC TTGACGATTG
	TITICAGITG CGGCCTITCT TCTCGTTCTT TAATTTGAAA CCTAGATACA TGCAGTAAAA	180	ACTIVACCAT AGRICUITAA ARAACTCICC ACACCITITI CITAGITIAE CICERCANGE
5	ACTAGGAGAA TGACTTTTAC CCTTGGGGAC AGCCAAGTTT TGTTGATAAA CCTATTTCCT	240 20	AGGGIGTICA GCACCIGIT CAAAGTCATA TITITICHGGGA AARATITICCA GCAGGTGTTATA
ì	AGCATGCCTT CAGGAAGTTG TGCCAGACCC TAGATTGTGA AGGACCCACT GTTCTTCTGT	300	GOACHTHAGC COACHCHGIG BACCCTHARTH TOTHTHABAC TICACCARTHABA THORARAMAN
	TOTACGAGOT COCTGAACCA TIGITICAGAG GACCAATGIC ACATCIOCITIC ATGGGGATGG	960	ACTIVALABITOR ACCOUNTS AND ACTIVATION ASSESSMENT ASSESS
25	NCATOGGAG CATCTGGGTG ATAXCTGTCT ACAGTATTGG CTCTTCTGGG AGGCTGATAC	420 25	ANTONIO PROCESSION OF THE PROC
	ACANGSCOTO TOTTCCACAT GATCATITIGG AAACCTCCCC CAGCCCCTAG CATCCAATGT	480	
Ş	GENAGGNAN CANGARCTOC CTGNAGNGA GTCCANGCTA CAGATACACA GCGTGTGCAT	540	(2) INFORMATION FOR SEQ ID NO: 22:
3	TOCOGCTOTC ACCITICATE TOCCACTICT GIATICTICAG AGATOCTICG TOGATOTITIC	009	(1) SEQUENCE CHARACTERISTICS:
	CITAACCICA GCIGACTICC CIGIGAATGI CIAATGCIAG ITCAGGGCCI CCAGGCATTG	099	-
35	ATTIVITACAG IOSTINACTIC CANTGAGSCT TCIOTINICA ITTIVSTOTICC TITIVICIGIC	35	
	attaaagaa atgattttcc c	741	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:
ç		04	CCACGCOTCC GGAATTCCCC TGAGGATCTT GGGCTATCTT TGACAGGGA TTCTTGCAAG
⊋		P	ITGATGCTIT CTACAAGTGA ATATAGTCAG TCCCCAAAGA TGGAGAGCTI GAGTICTCAC
	(2) INFORMATION FOR SEQ ID NO: 21:		AGAATTGATO AAGATGGAGA AAACACACAG ATTGAGGATA CGGAACCCAT GTCTCCAGTT
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 991 base pairs	45	CICAMITICIA AMITIGITICC TOCTGANAMI GNINGTATICC TGATGANTCC AGCACAGGA
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double		GOTCHACTAC ANCTCHOTCA GARTGATCAC ANANCAAAGG CAGATGATAC AGACACCAGG
	(D) TOPOLOGY: linear	\$	GATGACATTA GTATTITIAGC CACTGGTTGC AAGGCCAGAG AAGAAACGGT AGCAGAAGAA
ဝ	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 21:	200	GITIGIATIG AICICACTIG IGAITOGGGG AGICAGGCAG TICCGICACC AGCIACICGA
	GGCACGAGTC TCCCCTGGGG AAGTTTTTCT TTTTCAGGAG GGAGGAGGG TTTCCCAGGT	09	TCTGAGGCAC TITCTAGTOT GTTAGATCAG GAGGAAGCTA TGGAAATTAA AGAACACCAT
55	AAITOTOTOTA GAGTOTTGGG CAGAAAATOT GGGACCACAC CACACCAGTT CTCTCCTTAA	120 55	CCAGAGGAGG GGTCTTCAGG GTCTGAGGTG GAAGAAATCC CTGAGACACC TTGTGAAAGT
	TCCACOTCAT TIGCCTICIA TCCCACIAT GITTCCAGIG TCCTCTGGGT GITTCCAAGA	180	CAAGGAGAG AACTCAAAGA AGAAAATATG GAGAGTOTTC COTTGCACCT TTCTCTGACT
	gcaacaagaa atgaataaat ctetggtgag ttgtttattt gttettcaet ttgttttaca	240	GAAACTCAGT CCCAAGGGTT GTGTCTTCGG AGGCATCCAA AAAAAAAAA AAA
.0	CICTATITIC IGACHITAIG GGIGTCIGIG AATTAAAAAG GAAAAGIAGA AATBAGIAAA	09	

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55 50 5 6  $\frac{3}{2}$ ၓ 25 20 2 5 8 TOGATOCCCT GACTGAGCTG CITAGCACTG CCCTGGGGCC GAGGCAGCTT CCAGAACCCA TGGATATCCA GTATACCAAA GAGATGAAGG GCTGGGCACT GGTGCTGGCA SGARCTGGCA TTGGACTCAT GGTGCTGCAT GCAGAGATGC CCACTCTCCA GCTGGCTAAA GTGGGKAGGC CTTGGCCCAC CTGAGGCCCC AGGTGGGAAC ATGGTCACCC GOTGOTGAAC ATCOTCTCTG CCACTCCTGA CCCAGCCCTG AACAAAGCAC CTCAAGTGCA GCCAGCAGTC CAAGTAGCTG GACCCACGAG GAGGAACCAG GCTACTTTCC CCAGTACTGA CCGTGGTGGC CCGGAAGCTG GAGTTTAACA AGGCAGAGAA GCACGTGCAC AACTTCATGA GCAAGATCGT YTGCCTGTGC ACTGGAGTCA TGGGTGTCTG CTGCACAGCC CTGCTGGTGG TITGCCTIGGG GCCCTTGAGA CGCCGAAAGC GCTTGCTGGA GCAGGAGAAG TCTCTRGCCG CCTOGOCAGO AAGCTGGCTG ACCCCCAAGA CCCCGGGGGC CATGGGCGGG GATCTGGTGC GOCAGOCTICA CGACCTOCAA GCCACAGTOG CTGCCCTIGTG CGTGCTGCGA GOTGGGGGA AGGACCAAAG GOGGCCCTOG CTTOGAGTGG GTTGGCTTGC TGATGGCTGC TGGAGGGGAC AGAATOTGAG CAGOTCACAO COGGOOCTIGG AGAAACAGAT TGACACGOTG GOGGGGAAGO AACAAGTGAA CTCCATGGTG GACATCTCCA AGATGCACAT GATCCTGTAT GACCTGCAGG ANCTOCTOGO COCCATOAAC OCOTTOCOGO AGGTGCOGOT GAAACACOGG AAGGTCCOGG GGATOTICTA CAAACATACT CGCAGGAAGG AGICTCAIGC IGCCCGCANG CATCAGCGCA TGATICCCCAT CACATTCCTG ACCATCGGCT ATGGTGACGT GGTGCCGGGC ACCATGTGGG TOTOGITICOG GOGGIOCICO OCTOTICAATO CCACTOGGCA CCTTTICAGAC ACACTITIGOC 2 INFORMATION FOR SEQ ID NO: 23: E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: TACCCTCATC AAAAACACTC TCACTATGCT GCTATGGACG GATICCAGGAT TOTGGGAGGC TTCAGTTACC GOTGGCCGAG CTGAAGAACT AGTOCAGGCG ACTOGAGGCA GGACTCCTGG GTCCCTGGGA AAGAGGGTAC SEQUENCE CHARACTERISTICS: AGAGCTOCAG AGAGCACCTG GTGGGGAGGA AGAAGTGTAA CTCACCAGCC CTGGGGGGG GCTGGAGGTG GCGCCCCCTG GTGGGACAAC AAAGAGGACA (A) LENGTH: 1486 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear AGTOCGCTGC CCGAGTGCTA CAAGAAGCCT 23: ACCTCCAGCT 1140 1440 1380 1320 1260 1200 1080 1020 960 900 840 780 720 660 600 540 480 420 300 180 120

8 SS 8 45 ઝ 30 25 20 5 TATATTTTCC AGCATGTCTT GACAACCTGT ACTCTTCCAA TGTCATTTAT CAGTTGTAAA TACCITITICT ATTITITATIGA TYAGATICCIT TOTATTAAAT TOCTTCTCAA TGATICCATTT CAGAATICCCC CAGATGGCTT CATGGCCCCCC AAAGCATGGA AGGTCCTGAC AGATTATTAC TIGICICCA TOCCATOCCI GITAATICCI GGAIGGGGAC AAIGICCGIC AIGGCCITAA GICTAATAAG AGAAGICTIA AIGGCCICIG IGAATAAIGI AACICCAGIT CTANAAGAGG TOTOGCTCAC ATCAAGATTC GOTCCCAGAT TICTIAAGGC TITGITTICAC CATGIGICIA GITACITGCT GAAAAGIGAA TICTIATACA TITCATAATA AAATTAGCTC TATGTATITT CTACTGCACC TGAGCAGGCA ACACACAT ATACATACAA AGTCAAACTG AAGACCAAAT CTTAGCAGGT AAAAGCAATA TAATCITITA TAATGAAGTA AAAGTTGTGT CTATAATTAA AAAAATATAT ATATATATAC AGGICCCIGC AGAAGAACTA AGCCITIGGT CCAGAGITTIC TITICIGAAGI GCICITIGAT GUAGGAATAA GAGAGGCAA GTOOTTOGAA CAAGGGGTGG GTTCCGAGGA TGTACCGTGT CONTRAGON CICCOICOG ANCGANGIGO TOCCOCOGCO GCCGCCGCCG TCCCGCGICC CITICOCCOTT TETECTOCCA GOGGAGGTCC COGCITICCCG TOGAGGCTCC GGACCAAGCC GCCTTAATGT TGTAATCATA TCTTACGTGT ATATATCAGA TOTOTCCTCT TCTGTACAAT TGACAAAAAA AAAAATTTTT GOCTAACAGG TOTOTOOT GOTGGGAAAA ACAACGATAA GTTTTGCCCT GGAGGAGTAC AGCAAAAGAC GGAGAACCAG CAGAAATCCA CCAATGTAGT CTATCAGGCC CACCATGTGA TICOGICICI OCICOCOGOA COCOGCICOS COCAGOCAGO CAGOATOTOS GOGATICAAGA AATAACOIGA ATTITATCIT AGAGATCIGT (xi) SEQUENCE DESCRIPTION: SEQ ID NO: (D) TOPOLOGY: linear TGAAGACCTG ACTGGAGAAA CAAAATGTGC GCAGCCTATT TCTGTCACAA TICCIGATAT TITACCICAT GCTGTACAAA ACACGGTGAC AAGTTATATT TITICICACI 1200 1140 1080 1020 960 900 840 480 360 780 720 660 90 540 420 300 240 180 120 8

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TITTAATAGC ATACAGTGAT TIGATGAAAG GACGICAAAC AATGIGGCGA TGICGTGGAA

AGITATOTIT COOCTOITT GOTGTGGTCA TIGTGTCTTG CAGAAAGGAT GGCCCTGATG

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INFORMATION FOR SEQ ID NO:

24:

(i) SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: doub) (A) LENGTH: 2323 base pairs STRANDEDNESS: double

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AACTCGAGGG GGGCCCRKAC CCAATCWCCC TATAGTAKAC GTANNN

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CAAAGTTCCT

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TCAAGCTGTT

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CAGGCCTTTC TOSTCCTGAT AGGSTGGAGC AAAAGTGGAA AGGAAAGGAA AGAGGCTTTT CTCACAGCCA TTATATTAAA TAGTAGGTCG ATTCACATCT CGTGCTCCTG GCCACCTTCC CCTGTGCCTC AGTGACATGT AGATGACTGA CTGCCAATAC TTGTCACCAT TCCCTGGAAG CAGCTACCTA GOGGAAACAA GATGTAGTGC TATTGCCGAT AACAAGTAAG ATTTTCCACA CTGAGAAAGG AAAGCAITCG GAICTGCTGC AAAAACACAT ATATCCATAA AGACTCAIGT PATTCAGAAA ACAGATTGTG AACACAATCA CATTCGCATG AATCCTTTAA AAGGAAGAAG SCHTATCIGA HIATTAGAGA TATTATTITIG GAFATGITIAC TRATTAACITI OCHATGGCTG GCTTCTTACC AGAAACAGTA GGAAGAAACA CATGAACTGT GTACAAGACA TGAAACATTG ACCITAAAGT ATCTGCAAAT CTGAATITICT ATTTATTCCT TCACTGAATA TAGAAACAAT STANCCATGA TAAAGTCTGT TAITAATAAC AACATAATTC TTTTTTTAAA GAAGAAAAGC TRATITITICA TICACAGIGT ATAGATITTAT CTACITAGITI GIGITITIGCT AITAGIGITI DARTITITIT TITAAGITGA GIGITIGATA AATTITAAGA CCCIGICCCC ACCTIGITITI GAGTOCTISTIS TTGACTACAG STATATAGCY CAMTITAAAA ATCCTAAAGC AAAAGAATTT INITIATAAA AGAATCMAMC MOTTGCATGC ATGAGGCTGT GAAGTCAGAT ATTTAGTAAT AAAACCAGCA GIGCCIITITI IIGIAITITAC CCAITGACCC CCACCAAAIG CAACIGITITI ATATTAAGAA AATAGTAACA ATTITTAAAAT CTCAGAGTAA AATCTATTTC ACTACATGCT ITTCCCCCCT TGTTCTGAIT TAAGCAGTGT GTACTTGGCA TCTCTACATT GTCCTAGGGA CAGIOGICIT CIACAAIAIT AICAIGIAIG AIGITITIAIT GGICCITITIT AITCAIAGIG CTGCTGATAT GITGITITIT CACATGCTIT TGAGITITICA CITITITAAAC GAGAGCCAGC AAGCAAAATA GATGTGGCTG GGTCTGCCTG TCCGGGGGGC TYTTTGCACC GAGCTCTCAA ATCCTOTOTOTA ITGAGGOTIC CITITITICSTA CICAGGATIG GAGCTACAGC TGGGCCCCCC TCTCTCCCAT TCGTTTGAAG AGACACTGAG GGAAACAAGG GTTTCTTTTG AGGTGTCCTT **3**6: SEQUENCE DESCRIPTION: SEQ ID NO: 254 (A) LENGTH: 2036 base pairs STRANDEDNESS: double TYPE: nucleic acid (i) SEQUENCE CHARACTERISTICS: TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 26: CTAAAAAAA AAAAAAAAA AAA <u>@</u> <u>0</u> 0 (xi WO 98/39448 2 2 8 23 ಜ 35 6 5 S 55 8 PCT/US98/04493 1380 1440 1500 1560 1620 1680 1740 1800 1860 1920 1980 2040 2100 2160 2220 2280 2323 ŝ 120 贸 240 8 360 420 CAGCAGCAGC GCCAGCTGTA ATAAAAATA ATTCACACTA TCAGACTAGC AAGGCACTAG AACTIGGAAAA GACCACAGAA AACAAAGAAT CCAACCCTTT CATCTTACAG GTGAACAAAC TOTGATGATG CACATGTATG TOTTTTGTAA GCTGTGAGCA CCGTAACAAA ATGTAAATTT AGCTGCTTAA AGCTGCCTTC TCGGATACTG AAAGGTCGAG ACTOCCAGIO TGAAAAAAA AAAAAGCTAT TCCAAAGATT GAGAGGCAAT GTTTTTACTT CAGCACCTIT TAATATATAG TCTTAAAAAA CACTAATCTT AACTAACAAA AGTTCTTTTG AGAATAAGTT ACACACAATG CICTCTCG AAGAGACCTA AATTAGAAAG AGAAAACTGT GACAATTTTC ATATTCTCAT GCCACAGCAG TITIGICITTA ATAGTATAGT GCCTATACTC ATGTAATCGG TTACTCACTA CIGCCITITAA AAAAAAAAC CAGCATATIT ATIIGAAAACA IGAGACAGGA ITIATAGIIGCC TTAACCGATA TATTTTGTGA CTTAAAAAT ACATTTAAAA CTGCTCTTCT GCTCTAGTAC CATGCTTAGT GCAAATGATT ATTTCTATGT ACAACTGATG CTTGTTCTTA TTTTAATAAA GOCACGAGGC TGTGTGGTCA TGTTCCTCGT GGTGCAGTAC CTGACATGAG CCAGCCACGC TCAGTGGCTG AACAGCATTC CCACAGCCTG CAAGTGTGTG TGTGTGTGAA AGAGAAGAG GGCCCAGAG CCCCTTTTG AATGTTTGC CTGTCTGAAC TGTGAAGACA CTTGGGAGTG CACACCATGC GCGGTGCTTA GAAATGAAAA AGTCCCGGGT CTGTCTCTCT CACTCTCGCT CTCATOGGGG AGGGAAAGAA TGGCTTTGGT GGCTTTGTTC ACACAGCTGA TGCGTGCTGG GAAGGIGICC ACAGIGAGCC IGTOTGCAGG ACTOTGCACA COGITCACAC ITGICACAT ATTGTGGTCT AATTTCCAAC CTGCTCTGTT TYCTGTGACA TCTTGGAGGG GAGCTAGTGC GCCATTATTA GGAAGTGCTG GTGGCAGTGA AGAAGCACCC AGGCCACTTG TESTACCETS TETACACEAS ACAACACAGS AGCTGGGTCA GATTCCCCTC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25: 23 TATGCATAAT TCATTGTTGC CAAGGAATAA AGTGAAGAAA TITACITICCT (A) LENGTH: 683 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 25: CGAACAGAAA GTGCTTACAA TITITCIGAAC TGCACTGATT TTATTGCAGT CTGAGACATC TTCTGATGGC

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GOGCTOTOGO TETETOTOGO CETOGRIPAG AGGERICACE ATTRICTOGA ARERIGEAGT	CYCTTIACCT TIMICATTIA GGTGAGGCAT TOCACAAAAA CTCTCGACTT TGCCATATAA	CATTACTOCT CTACCCACCC ACTTICAGCT CCCCAAATTA ACTAGTOCAG TIGACTAATC	TOCCAGCTIC GGGAATGACA CTGCAAAGCT GATGCCAGAA ACTGCCAGAG TAATTCTCCT	CTCAGAGGCA GAGCCCAGCA CAAAACAGCA ATGCTAGAAA GTTACAATTG GAAAGTTTCC	ATTOCTOGAG AGAGCAGACT COCACCAACA TTCAACCCCA GCGCTGATAT GACAGTAATC	AACATTGAGA TGACATTTCC ATTTGAGAAG AAAATAGTTG CTTTCAGTGC CTTTTATTTG	GOCACGAGAT AACATAGGCA CAATAATACT GTATGTCTAC TTCTAGGATT ATAAGGAATT	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 27:	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 717 base pairs (B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear	(2) INFORMATION FOR SEQ ID NO: 27:			GAAACTGAAG GGAAATGATT AATACAAGGT TTTGTAACAA AAAAAAANAA AANAAAA	TIMPICATITI CCICCCATGI AACIAAGAAT CATGOCIATA TITCATATCA ACGITATATT	TTTCAAANG TIGTTACCAG TGAAACACCC TIGTOGITTA AACTIGCIAC AANGIATITA	AGCTTTACTG ATATACAGAT ATACTAATGT TTGAAGATGC TOTTCTTTOC AAGTGTACAG	AAGCAATGCA TATTTTTTAA ATTTGTCATA TATTGAAAAGA GCATGTTTGT TACATGTAAA	сстотттес тотагаласт тасполасал адалататт теоссствот техноттосс	COTOCOTIAT TRANSPARA IUTIUANDIA ITERRIOGRI ANDORAGIA DOCTITICOT ARCTIVICTI TITOTIGRATO COCTOTOCA GCATCITAT TOTRARIOCA TOCCITICOT	TARTICCTCIG GGAGGAATAG GAAGAAAACA GGAATGTTAA TAATGTCGAA CAGAAAACTT	GITTGCAAAG IGAGITITAT TITTITIGINA TITCTITAT TITMETINA WELWANDOO	THE SECTION OF THE PROPERTY OF	CACAMOGAGO TOTGATAACA CTAATOTTGA TITTITITT TITTACAAGI CATCAGRGAT	COURTER ACCUPANTAGE CHEMICALINATE TRACABAGAB ACCUPACAGAB GGATAATTIG	SONVERED SECONDARY SUCCESSION DESCRIPTION DESCRIPTION SECUNDARY	сторостваа тослаласка тосласская асосоросла ородновала асклюеств	остистоссь отосилост отидитотти тессемови осстовостк овозоствия	GOCTOCCTTT TACGOGARGO GAOCCTTCTC COGARCTTTT GIVETICIOC ACCICTIGIA
GOCATCACC ATTAICTGGA AACATGCAGT	IOCACAAAAA CTCTCGACTT TGCCATATAA	CCCAAATTA ACTAGIGCAG TIGACTAAIC	MIGCCAGAA ACIGCCAGAG TAATICICCI	ITGCTAGAAA GITTACAATTG GAAAGTTTCC	TCAACCCCA GCGCTGATAT GACAGTAATC	AAATAGTTG CTTTCAGTGC CTTTTATTTG	TATGTCTAC TTCTAGGATT ATAAGGAATT	SEQ ID NO: 27:	S: paire id				ITGTAACAA AAAAAAANAA ANNAAA	ANGGCIATA TITICATANCA ACGITATATT	IGIGGITIA AACITIGCIAC AAIGIATITA	IGAAGATISC TISTICTITISC AAGTOTACAG	anggaaaga gcangtingt tacangtaaa	HANATATTT TIGOCCTAGT TCATGITGCC	CATCHOTAT TOTACATOCA TOCCTTTCGT	SAATGITAA TAATGICGAA CAGAAAACTI	CCITIAL ITIACIIMM GSIGMIGIG	מכייויייייייייייייייייייייייייייייייייי	TITITITT TITTACAAGT CATCAGRGAT	CANGARGA ACCUCAGAA GGATAATTUG	CIGGGAAAGC	заставаела отогологалал ласловеств	гсссидая асставастк авааастама	SCAPCITY GIVETYCIGC ACCICITOTA
480 60	420	360	300	240	180	120	60	45	40		35			1980 30	1920	1860 25	1800	1740	1680 20	1560	1500	1500	1440		1320	1260 5 .		1140
CAGITAGOTG GACATCACGT GGACCCAACA CACGCATTTC CTGGGTTACT TACCAAGGAG	GOCACTOSTO CITICITCIOT GCCIACTOST AGGGSTOCAG CAGAGIGGTI CAGICIGGGA	GACACCTCTT AGAAGAGCTG CTAGAAAGGC AGACAGCACC AAGCGCTTAA ATGAGATGGG	ACCTINACCT CATCATTCAT THOOGGAING CANOGCAANA CCATGATGAG AATGCCCCTA	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:	(D) TOPOLOGY: Ilnear		(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 556 base pairs	(2) INFORMATION FOR SEQ ID NO: 29:	CACAGAAATC CTAGG	ATGACTCAGT GGTGGCACAC TATGGAGTCC TGCCCACAAG TAGCACACAT CAACCCACTA	AKGCTTCGAA GARGTAATAR AMCCCTGGAG AGAGAAACTG AGACATGTAA GAGGGTGGGA	TIMITITITA AMANGCIOC ICCAGGAAAT GCATATMAGG GCTAATCACC CAGIMITITO	ттепозосал ттемсестта самсатоско тооссетаса самалласто самстамала	AGGICATITO CTOCTACIAG CTGAAAGICA CCCCTGGGGG GCTGGGGGCCC TCGTGGGAAGT	AACTGAACTG TIGITTITCAT AGGTAAATGA GAGACTGAGT TITTICATIT CIGAAGAGAA	TATTOCTITT TOTOCACAA TTAATOTTGA TICTOCCTOF CIGTOCACAT TIGGATGAGG	GAATTOOOCA CGASCASCAT CCTAATTITA STITISGAGAT SCATICTAAA SGATCTICTC	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	(A) LENGTH: 495 base pairs (B) TYPE: nucleic acid	(1) SEQUENCE CHARACTERISTICS:	(2) INFORMATION FOR SEQ ID NO: 28:		GTCAGITGAT CAGACATIAG ATTATITATI OCTAMANCIA ADDRASSIA DESPESO	ומנומנות מודינות מודינות וודינ	GATTAGATAG GGATOGTOGC GTATCTTCCT ACAGTTTCCC TGTTAACAAG AAAGTCAGAG	Gotgatictca tacatisctaa agtitisasaa ccattisagta aagttaatisc ättaagaaga	AAATGCAGAT TCTTCATCTT CTCCCCAGAC CTCCTGAGTT AGAAATTCAC AAGTTCTCCA
TACT TACCAAGGAG	GIGGIT CAGICIGGA	CTTAA ATGAGATGGG	ATGAG AATGCCCCTA							AT CAACCCACTA	TAA GAGGGTGGGA	ACC CAGIAITITIG	TIG CAACTAAAAA	CC TOGIGGGAGT	TT CTGAAGAGAA	AT TIGCATGAGG	AA GGATCTTCTC							TO COMMON	TA ADADADA	AG AAAGTCAGAG	oc attaagaaga	C AAGITICICCA

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;	535		258	}
	ANTHONANGC AGGCAGATET TTACACCAGC TETTACETON TTGCAAACA ATGGAAATGC	300	CTGGCCTTAT TGGACTCCTT TTGGCTAGAG GTTCAAAAAT AAAGAAGCTA GTGTATCCGC	Ŋ Ŋ
v	CCACATOTICC ACAAACAAOT KTOTIGOTICTG CCTOTIGCCAT GAAGCACAGT GTGGCTGAGC	360	CHOSTITICAL GGGATTAGCT GCCTCCTCT AFTATCCACA ACAAGCCATC GTGTTTGCCC	g
n	OTCANGAGTC CCCACACTCA AAGGAGGCAG CAGATACAGG GCTGCACACT GTGTGATTCC	420	AGSTCACTUG GEAGAGATTA TATGACTUGG GTTTACGAGG ATATATAGTC ATAGAAGATT	Ę
	ACACATOTICA CATTICTUGAC ACGGACATOC TOGATGGCAA AACGAGCATC GGGCTGAGAG	480	TOTGGAAGGA GAACTITCAA AAGCCAGGAA ATGTGAAGAA TTCACCTGGA ACTAAGTAGA	5
01	GACTICCTGAG AAGGGAAGG GGCCTICCTGG GATGTGGGTT GATTGTAGCA GTAGCTCATG	540	10 AAACTCCATG CTCTGCCATC TTAATCAGTT ATAGGTAAAC ATTGGAACTC CATAGAATAA	ş
	GAGNISTGAC CTCAAA	556	ATCAGTATTT CTACAGAAAA ATGGCATAGA AGTCAGTATT GAATGTATTA AATTGGCTTT	E
;		<u>v</u> -	CTICITICAGG AAAAACTAGA CCAGACCTCT GTTAICTTCT GTGAAATCAT CCTACAAGCA	 ජ -
15		•	AACTAACCTG GAATCCCTTC ACCTAGAGAT AATGTACAAG CCTTAGAACT CCTCATTCTC	
	(2) INFORMATION FOR SEQ ID NO: 30:		ATCITICCTAT TTATGTACCT AATTAAAACC CAAGTTAAAA AAAAAAAAAA	
1	(i) SEQUENCE CHARACTERISTICS:	06		
20	(A) LENGTH: 434 base pairs (B) TYPE: nucleic acid	Ň		
	(C) STRANDEDMESS; double (D) TOPOLOGY: linear		(2) INFORMATION FOR SEQ ID NO: 32:	
;		36	(7)	
22	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:	4	(1) SEQUENC (A)	
	CTAAATGGTG ACTGTGGCTT TGTGGAGACA GGCCCCAAAT GGTAGGTGTG AACACAACAT	09	(B) TYPE: nucleic acid (C) STRANDENESS: double	•
ç	SCACAGAATG AGGAGACATG CAGAGTGCTG AAATACTGTC CTGGACAGAT GTGTTACATG	30	(Q)	
3	ACTITICITIT CAGCITALIT CTGTGGCCTG CCTTTGAAGA TAGAGCTTTG TTGALALITA	180	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 32:	
	CATTAAACCA AMITGIATAA YIMIGIITCCA TICTGACATG TIMITIAGCA AARGAAAAR	240	GAGCAGTIGC COCCAAAAG GAACCTTCCT CTACTTCCTG CCACAGACCC TGTCCCCACA	ส.
35	GAGTAATTET ACATCAGCAT CTTTAGTGCA TGCTAAAAGA TTAAAAATGT CTTTTGGGGA	35	CACTIVETIGE COCTIGETETS CTGGGAGGCE ACTIVECTECE CCAGTGCTGG ATTICEACECE	 я
	ACATGITITO TATACATAAA TGITTAGATA GAAATATTTA TAGAATACTC TATGIGAGTA	360	CAGCTCACCC TCAAACAIGG COCCCTCTCT CCTCCTGCTT GCCCCTCTCT GCTCCCTGGA	
,	THUNICTICC TRIGIAIRT TRIBICTAGA TOTOTICARIC TRIGIATICA TRIGIAARIOC	420	GGCTGTTCTG TCCTCCCCTC TTGAAAAGCA ATGCCAGCTT CCTGGGATCT TCTGCCAACT	F:
9	tatgatagt gaga	434	CCAGCTACCA TGCCCTTTGC TCCTGTCAGC TCAGCTCCTC AAGGGAATTG TCTAMCCTCG	 
			GIGICATIGAT TECCTOCATE AACCTCATCA CECTGATCA AGCTGGCATE TECCECTCA	 
. 45		45	CTOCACAGAA COGNICCCCC ACCACCTOCC TITIACAGGAA GGAAGAAGA ACATGGAAGA	
	(2) INFORMATION FOR SEQ ID NO: 31:		ANGGAACTAT AGGGGTACA ANGATGCTCA GCTCTGATCC CGAGGCAAA AAGNATCTTT	F-
20	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 715 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	50	ממסונים	
ý	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:	55	(2) INFORMATION FOR SEQ ID NO: 33:	
C	CCACGCCTCC GATCTCACAG CTCCGACACT ATTCCGAGCC ATACACAACC TGGTGTCAGG	09	(1) SEQUENCE CHARACTERISTICS:	<del></del>
	AAACGTACTC CCAAACTAAG CCCAAGATGC AAAGTTTGGT TCAATGGGGG TTAGACAGCT		(B) TYPE: nucleic acid	
9		180	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	
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	437	420	300 360	240	180	120	60								725	720	660	600	540	480	420	360	300	240	180	120	
60		55		50		45			40		Ę	36		30			25		20		ì	15		10	•		
granectang renesconce assussings thineaangn nethograan thinitiena $oldsymbol{0}$	GGCACGAGAA ATCTTCATGC TGTAGTCACT CCAGACCATG GAGTGGCTTT CCAGCTGAAT	5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:		(1) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 604 base pairs  (B) TYPE: nucleic acid	(2) INFORMATION FOR SEQ ID NO: 36:	5		калалала алалалала алалалала алалалала ала	) agtgantigt atgtggttaa ttataaataa aactggtacc aggnaaaaa aaaaaaaaan	TTTGGTTTTG TGATATTTNT GTGNATTAAG GNATAGATGT TAACCUTTAT TTTGTAGNAA	NOTCATTNIC AACCNITGNA TIAAAGCITA GACTAAANAG TAATATATNG TOGGNAGGAT	CAGATTACCC ATNTGCAGAA CTAAGGGAAG CNATTTATGT ATGAAAGNAA TTNTTGAATT	actemptict transcett tightcinin aggitrium atgacagicc inatetcama	) TOCACTIMAT ACOCACCIAT TINICAMBA TOTTATITITE TOGNIMOCAT TITTITIACC	TOTICCTITAG TOCCCIAAGG CTAAATTITIG GICATTITGAC ATCAGAGATG TIGTAAGTAT	TTAICHTCAC TITTHIGTGIA ATAAANTOTG CCATACTHGT ATGTGCTTCA AAAGGGCAAA	AMACAMINGI TITTITHAGC MITTGACING CICITIMAMI AGICIMCAMG ACATICACGI	TTGAGTGAAA TATCATAAGA TGANAATGGA AANNAGGAGA CACAAANAGT TATTNACAAAA	ACATTICITA CACCCIGGCA GAAGGAAGAG AAAIGTOITI 100GGIGGGI AACTAAAITI	TOTTIGITICA TENACATIOCE ANTECCICIT CAGIGITTAE TETCHAGIGA CAGAATOCIA	GARIGACCAG AIGCITAIGG ICIACATITI CCITTAICCI GITAGIATTA CCITCCITAA	AGAAGTACTT ACCTCTTGAA GATTTAATIAT ATAATGGTTG ACATGATIACA TGTACATGAT	GOCACGAGOT GGAACAGAGA CTAAATCCCA CGAAACTGAC ATTOTTAAAC ACACTAAAAC	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:	(D) TOPOLOGY: linear	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	
120	60							943	900	840	780	720	660	600	540	480	420	360	00E	240	180	120	. 60				

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(2) INFORMATION FOR SEQ ID NO: 34:

(1) SEQUENCE CHARACTERISTICS:
(A) LENOTH: 417 base pairs
(B) TYPE: nucleic acid
(C) STRANDENNESS: double
(D) TOPOLOGY: linear

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AAAAAAAAAA AACTCNA

CTACATCTGC AGCTTGAGGT TAGCCTCATA TCACATTACA TICTCACTAN AAACNAAAAA

ATCACCTGGC GTGCCCAGAT CCTCGCARGG CAACACCCTG TGATAATTCC AGGTGATTCT AGGAATCTTA TAACCTACGT GOACTCTTTC CATCCGTACA TIGICGIGCA CATGCCACTC

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CCAACGTGGA AAAGTATTCC AGGTCCATCC CCAAGGAACC AACACCGATG ACATGGACTC CICCIGCITT TATTOGCICT IGICGAAAIC AAATTGGAAG AICTICAGIC CCAGCIGCAC 3

TOCANGATAC TGAGAGATTO AAGCATGCTC TOGAAATGTT CCCAGAACAT TGCACGATGC

CACACAGCAT GCTGCCCTCA GACGTGTCCA TCCTGTACCA CATGAAAAACG CTGCTGCTCC

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:

30

AAAAA

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GTTCATACTT TICAATTIGA TAGAAATAAA GTTTTTTTCT GCTTATAAAA AAAAAAAAA TOTATATECT ATCCTAACTG TTAATTGTAT TATTGATTAT OTTGATTATC TTGCTTGAAG TIGGIGGCC ICITAATITI OGIGIAIGIG CIICCAAGIA ICIAAACCIC CAGICIGAIC 20

CAAGITITICCC AGAAGITOGIG TOTITIAIGAT GAGICAGAGI GCITITICCIC GGIOGGACAG

CAGTICTATA ACCCAATGAC AACCIGICIC TITIOGITTAC IGICCIGIGA AAIGICAGCI TCTCTGGGGA AATTGTTGAG TTACAATGGC ATTTCACTGT GATCCCTCTC AAGCTCAGAT

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TCTAGTTTTA TGAGAATTTG TACTACTGAT TTTTATATAT TCCTGTTTTT GATGAACAGA

5

AGTCAGAAAT GTCCTAAATA ACAAACTATT TTGTATTTAA TTTAGGGAAG ACTAAAGGGA

AGAAAAATGA AAACTCAGTC TITATGTAAG CTCCAAGGAT ATTAGGGCTT AAAGGGCTTT

S

GITCCICTGG TAATAATTAG GITATICCCA GAAGCACAGI GICATICTIT AAATAAAAGC

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

TITICCIOTITI AAAGCITTITC AAAGGAGCAG ACCACCITGA AGAITCCCCC TAGGGITGAT

ATGIGICTAA TICATITTAT AAAAATTAIT CIIGICTICA TITTAAAGCI TIGGCIATAI

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(2) INFORMATION FOR SEQ ID NO: 35:

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TIGGALCTIG GGAGTITICT TIGITIGCIC CTGTGTTTGC CCACCTITAA TAAAACCAGG 180	GGNACGGCAG CAGCGGTCAC AGGCNAGTNA ATMGTNANTGC CGGNACCNAGT	AGT TICCICCOCC	180
COCAMACAAA AACCATAGCA TTCTGAAGAA TAGGGGGCCC ACATTGGACC CAGTATGTCA 240	TITATICATES CACCCACTES GGTATATGCG TIGTEGTCTG CCAACTITGC	TGC CGTGAACAAT	240
	5 TTCAGCAATA ATCAGATGGG GGCTGGGGGA ATATTCAAGA TAAGGCCTGG	res chstrestecs	300
ACACATTTA TGCCATATAA TGGTGTGTT TCTTAATTTT GTTTCTTGTG GCGAAATGTG 360	GCTGATGGTT CAGTGCCTGC GSCACCGTTT YTGCCGTATG TTGCACACA	CCA GGNTCTTTAA	360
	ACAGITITICG SACCGCGITI AGGICAAGG GITCAANGCC GGICGGIAGC	AGC TOSTCCTTAG	420
	10 GITCACGGG AGCATAAGCA TIAAACATCT CATCAATITIG CITCTGGCTG	CTG GCGCTATCAA	480
TCAGTAGTAA TCTTGTTCAT GTGCTTTTTAC AGCCAGCTAC ATTTAAGGAT GTATTAGTTA 540	TACTITICCAG CATATIGITIA CGCTGGGGGA AACGGGTTAG CGTFTTGCCCC	OCC ARCINGWICAT	240
15 cagaaattat atgectests atstetet acteaataaa stacatecet ceacaaaaa 600	15 AGGCAATGGG CTTAATGAGA TAATGAAATA CACCACAAGG TAGGGCTTCA	ICA GACACCOTIT	009
AAAA 604	CCATATICACT GACTICCAGTA GTAAACACCA COTCCCCGGG ATAATGCGCC	SCC TOCACCAGIT	099
20	сагосмотал ат		672
(2) INFORMATION FOR SEQ ID NO: 37:			
(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 349 base pairs (B) TYPE: nucleic acid (C) STRANDECNESS: double (D) TOPOLOGY: linear	(2) INFORMATION FOR SEQ ID NO: 39: 25 (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1908 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double		
30 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 37:			
GTGAGTGCCC GGGAGCCCCG AGGCCCTGCC CCTAAGAAGG ATATCTYTRA CCGCTCCCTT 60	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 39:		
greatacce tracececa ectectease eagregeere attochasse ceteactess 120	AGAGITGATA TITITAGAA CAGTAATTIT ACTITIAAGG AAATTGOCTA GCTCTITIGAC	ста остетителе	09
35 GOCACATAGA GCATTTGGGG GACTGCGAGCTTT GACTTCCTGC AGTTCGGGGG 180	THINGAGCTG TAGGAAGCTC AACATTICTT TGTAGAGAAC GTTGCTTTTT	III IIGGAIIGIA	120
AAAACCAGAT CATGATGACC AAAGTYTAGA TATTCTTGAT CTTCATGGTG CTGATCCTGC 240	CAGGTATAAA AACATTGCTT TTGTTGAATT GTATAGGTGT AAAAAGGGAA	SAA TAACTOTATG	180
40 cenceetises retracease tribitisceae cacritecies retrabitica santaagaet 300	40 слосттал людалитот осттиосся талотсятля алтосситто	me tactterage	240
CACATCAGGA GAGCACTGTC CCCAGGANAA TGCAAACGGG TTGGCAGCA	CATTITATTI TCCTTTAGAA ATGGACATCA GCTCTTCTCT TCTGACTGGT	SCT AACACATAGC	300
	CCCAAAGCAT GAGAITAITIT TTCAITIGGT TITIAITIGIT GITTAGTITIT	TT GOTTIGITAC	360
45	4) GOCAGOOCAG TCTGTCTGCG GAACACTGAC TCTGCTCTCT AATGAGAACA	ACA AAGTTAGAAA	420
(2) INFORMATION FOR SEQ ID NO: 38:	TCTGCCGATA ACCTABABITA ATTITAGABAT GARITABABA TGTGABATCG GGTTABABGTG	TCG GGTTAAAGTG	480
(1) SEQUENCE CHARACTERISTICS: 50 (A) LENOTH: 672 base pairs	50 ATGATGATAA AATMGCATGC AAGAAACAAG CTCCTTCCAT CAGACTTGGC TACTGTTTTC	SGC TACTIGITATIC	240
	TICHGGING ATTIGGTING GARGAGCCIC INGITICCIT CICTITIGGG TAIGICITICS	soc ratererrec	009
(D) TOPOLOGY: linear	TITICITAATA TGITTICIAAC ATTAITICAGA TATAATICAC ATACCITACA ATTICACITAE	ACA ATTICACITIAL	099
55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38:	33 TITANGGGTA CARITTAGTG GTITITAGTG TAITCACAAA GTIGTGTAAC	AAC COTGACCACA	720
GTAGTEGTTG CGGTTGCCGG GATGCCGAAQ ATCTCGCCGT TTGAAGTCGT AAAACGCACC 60	STCANTITIA GACATITICS TIACCCCAAA AAGAACCCT STACCCTICA GCASTCACCT	TGA GCAGTCACCT	780
TOGGTACOGG TGCFTGTTGG TTTGGTGATT GTWATCGTTG CTACAGAGCT GATGGTGCCA 120	60 CTCATTITCT CCCAGTGCCC ACCCATCCC GGAGCCCCKG GAACCACTAA TCTATTICTC	IAA TCTATTTCTC	840

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	алалалала ала	СИПСАСТТІС СССІСЛАЛА АЛАЛЛАЛАЛА АЛАЛЛАЛАЛА АЛАЛАЛАЛА АЛАЛЛАЛ	CCCCCTTCTT TCTCCACTOT ACAGAAGAGC CACCACTOGG ATOGGGAATA AAGTTGAGAA	TITIAATCETT, GCTGACGOTT CAGTCCTGCC TCTACTGTCT CTCCATAGCC CTGGTGGGGT	TIOGRAGOGO CAGCCCCGC TOGCTTCTTO GETTITOTOGE TOCCAGCCTC AGGICATCCT	TIGGGAATIG CACTITIGGG CCTTIGGGCT CTGGAACCTG CTCTGGGTCA TIGGTGAGAC	стоссетств досогосств сапстетова достесттал товетесска десттоветт	АТСАСТВИТО СОСТОВОСАС СТСАОТСКСЕ АССАССАТВО ВОХОЗСТСКТ САЛАБАСАСС	AACGTGTTTG ACACAGGCTT GCGGGACGTG CAGCCCTACC TGTACAAGAT CTCCTTCCAG	спосысства теанпоска теантастте сосывается сенсасного сенсотоват	GAGCOCCAGA TCTOCCOGGA GAAGGTOGGT GAGAAACTCT GCGAGAAGAT CATCAACATC	TOCATICCCAT GGGAAGTGTG GACGGTCAAG GTGCATGTGG TAGCCCTGGC CACGGAGCAG	GOCCAGATOT CCTTGGAGTT CTACCAGAAG AAGAAGTCTC GCTGGCCATT CTCAGACGAG	ATCCOTOCTC CAAACTCTAC ACTCAAGGAT GCACTGCGCA ACTCTGGTGG CGATGGGCTG	стосовност сосносново сотсновнов навновные оснтаноска насттанова	TRACTIONAL TOTOGRACTIC CITETITOTIC CCTCAGCOCC CTCCCGAGGT TGAAGCOTAC	recharchae emagadada ageococce anancheeca nathaneera recretera	стотосское отважение сеозважена тоаннесска осостестве местисская	GOCTOGICCI GLACTOTOGI TOGGOGAGGI GGGAGCTOTI TIAACCGIGI GCCCCCCICIC	GOCACAGAGO CICCGACCCA GOTOOTCTOG AGCCTGCCGG GAGAGTGGTG GCATCTGAGA	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 41:	(1) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1153 base pairs  (B) TYPE: nucleic acid  (C) STRANDELNESS: double  (D) TOPOLOGY: linear	(2) INFORMATION FOR SEQ ID NO: 41:	CATAAAACCT CATATTITAA ATNAAGTTGA AATTIGAA	GENACICENT AGINTISTICS CITENCITICAS CITECRATISGS CANCITOCICE COCUTICAGA	CACCAGOCCC TOCCAGAACC TOCTCAGTTC CTTCACAGTG CAACCCTGTG TACTTGGCCC
	1153	1140	1080	1020	960	900	B40	780	720	660	600	540	480	420	360	300	240	180	120	60				458	420	360

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GCCATAAAGA GAAACAAAAG ACAATGATGG TATTCTCTGT GTCCTCAGCT TTGGCACTTT

TOTTGATGIT GCTAAGGAGC AGTGACCTTG CTAAAAAGAC TGAATAATCC ACCCACTGAA TAGCTAACCT GGGGAGGAAA TGAAAATTTC CTTTGTGGAT CTCCCCAAAT CCATTGTTGT

120 180 240 300

8

CCTCAAAAAA AAAAANGAAA GGAAAGAGGT CTCTACACAA GCCCGTGATT CTTCATGGCA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:

AGGGATAACA TCAGAAATOT TTCATTTYCK GCTATTAGTT TCCAITCCTT TCCCCATCCA

40

(2) INFORMATION FOR SEQ ID NO: 40:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 458 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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TGTTAAAAAA AAAAAAAAATT AAAAAAAACTG GGNGGGGGGC CCGGTACN

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TCTCAGTGGC TTGACAGCAT CTTCCTGGTT GTATGTGGCC TGTTTACATG ATGTATTGAA

CAGTCAATTA TIGCCTAGGG TAGTICAAAA ATAIGATGTG AGCTAGTTAA GCCTTIGCTT
GACTGATTTC AGTGATATTC AGAAGTGTGT ACCAATCAAG GCTCTTTAAA ATACGGAAGG
ACTCACTTAA TAACCAGGGA ACCAGCCAAA TACTGTGCAG CCGCAGAATA TGCATATCAA
TGAGTTGGAG GTGATTATTC TCTGTAACTC CCTAATGATT GTTTTCTAAG CATTGTGGCT

1560 1620

1680

TAATGITGIT TGITGIGAGC ATCAATGCCT GTAACACCAA ACTAAACACG TGITTITIGG

ATATOTTICC AATCITTAAA TGACCTIOCC CIVICCAATA AATAAATGAT TOTCTCACCC

1860

1800

25

20

GATGCGTTGT AACACTGCTA AATATGCTAA GTACAGAATT TTATCTACAG TACTGTGAGA

1320 1320 1380

1500

ACTOCAGIGA GITTOTOCIG TOTITITICAA GIATOTACCA TAGGACTIAA AGGIGATITIG

CAAAAATCAG GCCAAATGAC TIGGCAAATA ATIGACAAAG IGGITTICAC GIGIGICIAT CITTIGCTAG COCAAATTACT TITTIGAGTTIC

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TGAGAACTCT TATTOCTOTG AAAGGGAGTG GTTGGTAAAA TCAATAGATT TCAGGCAAGA
GGGCCAGATA CCTAACAGGT TTTTCTCCGT GAATCTTATG CTGAGTAGTT TTTCCTCATA
ACCAAGCATT TATGATATAT TACTACTTAT AATACTOTGG CTAGTCTCTA GAATGGAAGT

TGAAATCTTT GCCTCCTCAG TCGGGAAGAG TCCTGCTAAA AATCAGGCTA AAAATCAGGC

900 960 1020 1080 1140

S

TCTCTOTAGA TTTGCTTMTT CTOGTCATTT CANNAAATG GAATTCTACA ATMTTCOGTC TTTTGGGACT GGCTTCCCAA ATMTGATTTT CTMTMTGGAG TGAGAAAATT CTTCTCATCT

	TRACTICATICE TRAGTIGGECA CATTIAGACA TRACAGGOCT GATGACCAAC AAAGATGAAG  TRECETATICA ACGAAAGCA ATGACTICA GATACTCAAA AATGTETTCA CICTOTICTTA  AATTIGACAG TRATTAAAA GCTTAGTTTA AATGAAATTT GTATCCTTCA ATTITTTTTTT  10  CATTICACAC AAAGATTAT AAAGGTAAAG CICCTAGTTA ACTICATAAG GATTITTTAAG  ATACTCTATA AATGATTAAA ATTITTAGAA CITCCTAGTA GCAAACTGAA  15 TREATTGAGA AGGAAATTG TAACTCAGAT TIGTGATGC AGGAACATGA GCAAACTGAAA	266 TAACTCATCC TAAGTGGGCA CATTTAGACA TAGCAGGGCOT GATGACCAAC AAAGATGAAG TTCCCTATCT ACGGAAAGGC ATGACTGGGA GGCCCACAAG GACTCTCATT GAGTTCTTAC	1500
(2) INFORMATION FOR SEQ ID NO: 42:  (i) SEQUENCE CHARACTERISTICS:  (a) LENGTH: 1981 base pairs  (b) LENGTH: 1983 base pairs  (c) STRANDENESS: double  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:  GGCACCAGAG GGCCCAGCC GACAGATGT TCTTCCTGCC TCTTCCGCC GCGCGCAG  TAGTCCTCCG ACGTCTTGCC GTGAGACGTT TCGGGAGCCG GAGTCTCTCC ACGCCAGACA  TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTCCCACAGT  1		TAGACA TACCAGOOT GATGACCAAC AAAGATGAAG ICTGGGA GOCCCACAAG GACTCTCATT GAOTTCTTAC	1500
(1) IMPORMATION FOR SEQ ID NO: 42:  (i) SEQUENCE CHARACTERISTICS:  (a) LENGTH: 1983 base pairs  (b) TYPE: nucleic acid  (c) STRANDEDNESS: double  (d) TOPOLOGY: Linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:  GGCACGAGAG GGGCCGAGC GACAMGANGT TOTTGCTGCC TOTTCCGCT GCGGGCGAG  TAGTCGTCCG ACGTCTGGC GTGAGAGGTT TCGGGAACA AGAAGATGAT GTGCCACAGT  TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  10 TOTAL CAAAGAAAA AGAAGATGATT TCGGAAGAAAA AGAAGATGAT GTGCCACAGT  11 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  11 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  12 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  12 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  12 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  12 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  13 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  13 TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  13 TGACGAAGAAGAAAA AGAAGAAAAA AGAAGATGAT GTGCCACAGT  14 TGACGAAGAAGAAAAA AGAAGAAAAA AGAAGATGAT GTGCCACAGT  15 TGACGAAGAAGAAAAA AGAAGAAAAA AGAAGATGAT GTGCCACAGT  15 TGACGAAGAAGAAAAA AGAAGAAAAA AGAAGATGAT GTGCCACAGT  15 TGACGAAGAAGAAAAA AGAAGAAAAAAAA AGAAGATGAT GTGCCACAGAT  15 TGACGAAGAAGAAAAAA AGAAGAAAAAAAAAAAAAAAA		ICTIGGGA GACCCACAAG GACTCTCATT GAGTTCTTAC	
(i) SEQUENCE CHARACTERISTICS: (A) LENTH: 1983 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (C) STRANDEDNESS: double (D) TOPOLOGY: linear (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42: GGCACGAGAG GGCCGAGCC GACAGANGT TCTTGCTGCC TCTTCCGCCT GCGGGGGAG TAGTCGTCCG ACGTCTGCC GTGAGAGGTT TCGGGAGCCG GAGTCTCTCC ACGCCAGACA TGACGAAGGG CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GCCACAGTT			1560
(A) LENGTH: 1983 base pairs (B) TYPE: nucleic acid (C) STRANDEDRESS: double (D) TOPOLOGY: linear (Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:  GOCACGAGAG GGGCCGAGCATCT TCTTGCTGCC TCTTCCGGCT GCGGGGAG TAGTCGTCCG ACGTCTGCTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GGCCACAGT  TGACGAAGGC CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GGCCACAGT		ITCGITICAG ICAAGACAAT GCITAGITCA GATACICAAA AATGICITICA CICIGICITIA	1620
(c) STRANDENESS: double (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:  GGCACGAGAG GGCCCGAGC GACAAGATGT TCTTGCTGCC TCTTCCGGCT GCGGGGGAG  TAGTCGTCCG ACGTCTGCC GTGAGACGTT TCGGGAGCCG GAGTCTCTCC ACGCAGACA  TGACGAAGGG CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT  1		artiggacag tigaactiaa aagopititig aataaatiga tgaaaatctt tiaaggaga	1680
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:  GOCACCAGAGA GOSCCGAGC GACAGARTOT TOTTGCTGCC TOTTCCGCT GOCGCGAGC  TAGTCGTCCG ACGTCTGCC GTGAGAGCTT TCGGGAGCCG GAGTCTCTCC ACCGCAGACA  TGACGAAGGG COTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT		COACAC AATGAAATTT GTATGCCTTG ATTTTTTTTTT	1740
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:  GGCACGAGAG GGGCCGAGCC GACAAGATGT TCTTGCTGCC TCTTCCTGGCT GCGGGGCGAG  TAGTCGTCCG ACGTCTGCC GTGAGACGTT TCGGGAGCCC GAGTCTCTCC ACCCAGACA  TGACGAAGGG CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT		MUNICAL TOTAL STATEMENT SCHOOLS OF THE STATEMENT OF THE S	
GOCACAGAGA GOSCOCAGOC GACAAGATOT TCTTGCTGCC TCTTCCGGCT GCGGGGGAG  TAGTCGTCCG ACGTCTGGCC GTGAGAGGTT TCGGGAGCCG GAGTCTCTCC ACCGCAGACA  TGACGAAGGG CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT		STAND THAILTIE ACTION CALLITING	2
TAGTOGROCG ACOTOTOGCC GTGAGACOTT TCGGGAGCCG GAGTOTOTCC ACCCAGACA TGACGAAGGG CCTTGTTTTA GGAATCTATT CCAAAGAAAA AGAAGATGAT GTGCCACAGT		afactictata aatgattaaa attitttagaa cttcctaatc actittcaga gtatatgttt	1860
		ttcattgaga agcaaantig taactcagat ttgtgatgct aggaacatga gcaaactgaa	1920
	AATTACTATG CACTTGTCAG AAAC	aattactatg cacitgicag aaacaataaa tgcaacitgt tgtgcaaaa aaaaaaaaa	. 1980
TCACAAGTGC AGGAGAAT TTTGATAAAT TGTTAGCTGG AAAGCTGAGA GAGACTTTGA 240	20 AAA		1983
20 ACATATOTOG ACCACTOTO AAGSCAGSGA AGACTOGAAC CTTTTATGGT CTGCATCAGG 300	23		
ACTICCCCAG CGTGGTGCTA GTTGGCCTCG GCAAAAAAGGC AGCTGGAATTC GACGAACAGG 360		;	
25 маластовся тельновська вымасятся высстостот тосносовою тосновська 420	25 (2) INFORMATION FOR SEQ 1D NO: 49:		
TTCAMEACCT GENECTORS TOTOTOGRAPO TOGATOCOTO TOGAGAGOCT CAGGOTOCTO 480	(1) SUNCHLE CHARACTERSITES (A) LENGTH: 1406 base (b) more and a notal	E CHARACLEALINGS: LENGTH: 1406 base pairs	
COGRAGOGRACE GOTOCTTGGT CTCTATGART ACGATGACCT AAAGCAAAAA AAGAAGATGG 540		STRANGENESS: double	
30 CTOTOTOGOC AAACTICTAT GGAAGTOGGG ATCAGGAGGC CTGGCAGAAA GGAGTCCTGT 600	•	10) TOTAL TOTAL TOTAL TOTAL CONTRACT TO NO. 43.	
TTGCTTCTGG GCAGAACTTG GCACGCCAAT TGATGGAGAC GCCAGCCAAT GAGATGAGGC 660	AND THE PROPERTY CONTRADICATION	CONTRACTOR CONTRACTOR	9
35 сласскант тессалант атталалас атстелала тесплетнег лалассалос 720	35	AND CONTROL OF THE PROPERTY OF	130
TECATATEAG ACCEANGIET TGGATTGAGG AACAGGCAAT GGGATCATTE CTCAGTGTGG 780	TIND TOTAL TOTAL STATE OF THE S	ALMOSTUCERA RESELLCTAL RUSINILLORC ALLICLICAN COMPENSORA FANTELLORGICA. TRESINGUESES STOCKHESICS NUMBER TO SECTIONAL COMPESSORS PROFILES	Q Q
CCAAAGGATC TGACGAGCCC CCAGTCTTCT TGGAAATTCA CTACAAAGGC AGCCCCAATG 840		COTOTA SECULIARIO CONTRACTO CALLANDOS	
40 caaacgaace acceptedte tittofficea aaggaathae citteacaof gotsofahet 900	40 ATAAAAAATT CCAGTCAATT ATTC	ATAAAAATT CCAGTCAATT ATTCCTCAAC TUAAAGTTTA GTGGCAGCAC TICTATTGTC	240
ссательнос ттепесьмат апослестся телеросттем сатосластя 960	CCITICACITIA TCAGCATACT ATTIC	CCTICACTTA TCAGCATACT ATTGTAGAAA GTGTACAGCA TACTGACTCA ATTCTTAAGT	300
-	CIGATITICIG CAASTITITA TOOT	CIGATITIGIG CAAATITITA ICGIACITIT TAAATAGCCT ICTIACGIGC AATICIGAGI	360
CCCCTCTTTG TGAAAATATG CCCAGGGGA AGGCCAACAA GCCGGGGGAT GTTGTTAGAG 1080	TACAGGTAAA GCCCTGTTGT AAAA	TADAGGTAAA GCCCTGTTGT AAAATAAAGG CTCAAGCAAA ATTGTALAGT GATAGCAACT	420
CCAAAAACGG GAAGACCATC CAGGTTGATA ACACTGATGC TGAGGGGAGG CTCATACTGG 1140		TICCALALAO GALGITGAAA ALAGIAATGI GACIALALAO ITITITIAAL TGIAAGALA	085
50 CTGATGCGCT CTGTTACGCA CACAGGTTTA ACCGAAGAT CATCCTCAAT GCCGCCACCT 1200	JU TCACCIGSCT CITTAATATA TGA	TCAGCIGGCI CITTAATATA TGACTAAACA ATAATITTAAA ACAAATCATA GTAGCAGCAT	540
TAACAGGTGC, CATGGATGTA GCTTTGGGAT CAGGTGCCAC TGGGGTCTTT ACCAATTCAT 1260	ATTAAGGTT TCTAGTATGC TAAT	ATTAMOSCIT TCTAGTATISC TAATATCACC AGCAATGATC ITTGGCTTTT TGAITTATTT	009
55 cchasciete gaagaarte ticaagacca gcattgaaac agagaaccot gtetogaaga 1320	GCTAGATGTT TCCCCCTTGG AGTI	GCTAGATOTT TOCOCCTTOG AGTITITOTCA GTTTCACACT GTTTGCTGGC CCAGGTGTAC	. 099
TGCCTCTCTT CGAACATTAT ACAAGACAGG TTGTAGATTG CCAGCTTGCT GATGTTAACA 1380	TOTTIOIGGC CITIOITAAT ATCC	METHIGIOSE CHITOTIAAT ATCOCAAACE ATTOOTIOSG AGTEACATIG GTTTCTTAAA	720
ACATTOGANA ATACAGÂTET GEAGGAGEAT GTACAGETGE AGCATTCETG AAGAATTCG 1440	ANANANANA ANAGGRAT AGG	AAAAAAAAA AAACGACAT ACSTGACAGC TCACTTTTCA GTTCATTATA TGTACCGAGG GRACAACAACA GTTCCAAACAA GTTCAAAAAA CAACTAATTATA TTCATTTCTCA TCAAAAATAAA	780

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CANGARAGAC CTGCATGACA TCAGCACCAA AACAGGCATC ACCCTCAAGA GCTGCCGGAG ACTOCACCAG CTCATCTTCC AGATTCCGCC CTCCCGGCAG GCACTACTCA TCGAGAGGTA CATGGACCAT TACCGCACCT TCCACATGCT CGAGCGGCTG CTGCATGCGC CGCCCAAGCT GOTOCOCTOG GONATOCTOG AGCAGACTOG COCCACOGCA GCOGTOCTOC AGAGCGACAC COMPONED TATESCENCY GOLICOMING TYNCICOGIS ACCOMOSOG ISGECCISCS CCCGGGCGTC GCCATGACCA GTGAGCTGGA CATCTTCGTG GGGAACACGA CCCTTATCGA GOOCCIGAAG OCOOCREGEE AGIICCEGAGE AGIIGCIECGET CETIGEICOGG GEGETIGEGGE 2 GGCCGATTAA NTGGTTTGAA GNCTTG TAGITATGIA GITATTATGN AACCACCAAG ATTITTITTIGG CTATITACCG TAACCAAAGG TAAAATAGGA AATTATAAAT ATATAGTTTT AAGCCTGCAT CAGTGGGAGT CTTGGCTATG TCATAAGAAA TTAAAAGAAC TTACCAOGAA OGTTTTTTAAG TTAGAAATAT TCCATGCCAA TOTAMAGAAC TGAMAACMAT GTATATOTTG TAAMTATTTG TGTGTTGTGA GAMATTTTTTG CATGAGTCAC ACCITITIOIT CIGIOGIAAC CTATAAAAAA AGITTOTCIT TGAGATTCAA CTACAAGGAA TATTAAAAAA ATCTATTCAC TTTAACTTAT AATAGTTTAT GAAATAAAAA CTAATGAACT ACAGCTATCT TAATTTOGTT CTTCAAGTTT TCTGKTGCAC TTGTAAAATG TOGTOCTTAT TITITICAAAA ATTIGCIGIG AACAACGIGA IGACAACAAG CAACATITAT AAACCTIGIA TITAACICTI TICAATCCTI TIAGATAAAA IIGIICIITG CAAGAATGAT CCICITCITI CTATICCCTTT GATGAGGCCT TIGTTCGGGA GGTGCTGGGC AAGAAGCTGT CCAAAGGCAC CTCACAGATG GATGACATGG ACATGGACTT AGACAGGAAT TICICCAGGA CTIGAAGGAG GGACAATATT CAGCAACACT TCCTCCTCTC TGACCGGTTG GCCAGGGACT ATGCAGCCAT ACAGTITIGAC AACTITIAAAC GGGICTITCAA GGIGGIAGAG GAAATGCGGG GCITCCCTGGI COGNIGACINY OCCINCIOCG CIGAGCICAY GANCCAAAAC IOGACCCTIG GACCCOICGA INFORMATION FOR SEQ ID NO: 44: Ě GCTAACAACC GCTTTGAGAC AGGGAAGAAA AAACTGCAGT ATCTGAGCTT SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 44: (A) LENGTH: 1391 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 1260 1200 1080 1020 120 780 660 90 540 80 420 36 300 180 960 900 မွ 25 20 50 3 8 33 5 8 S 9 CCGTCGTGTG ATGCACTTTC AGCGGCAGAA GCTGATGGCT GTGACTGAAT ATATCCCCCC GOTECTEGET COSCIGLAGEA ACTOCICIOCO GTOCAGATOT OCCTOTOTET GTETETIGAGE OCCOTTETTE CTGAATCAGT ATTEAGEGTE TOTOCAATEC CTCGATGGET TECGACACCA GGGGGCTGGT GAACGTGCCG CCAAGCTGAC CCACAATAAA GATGTCAGAG ACCTGTTTTGT CICAMOGICC TAGIOOCIGA CAMOGACCIT CICGACCIGC ACAMGAGCCI GGIGIOCACI GEOGGEOGG CIGCOTACCO TOCAGACIGI COCCIATOGO TOCAAGGOIG TIACCOGGCA GECACGAGTG GAGATGOCTG CGGCCGTGGC GGGGATGCTG CGAGGGGGTC TCCTGCCCCA CIGAGOIGCC TCCCAACOIC COCCCACGCT GACAATAAAG TIGCICIGAG TITIGGAGACT GOCCETETGG GACCOCTACA TOGGCACCCT CCGCGGCTGC CTCCTGCGCC TGTATCATGA GGACCTCGTG GAGAAGTTTG TOGAACCCTG CCGCTCCGAC CACTGGCCAC TCAGCGACGT GCTCTCCCGG AAAGCTGGGC GTCTTCTCTG AGATGGAAGC CAACTTCAAG AACCTGTCCC GAGGATTICCA AGTACCAAAA TCTGCTGCCC CTTTTTTGTGG GGCACAACAT GCTGCTGGTC GANACCAGOC ATOCACOCAT CATOCCTOCC ATCTCCTCCC AGCCCCCCAC 3 CTCAGGGAAT GGTGTCTGTG CTCAGCCCAT CCACCAGAAG AGTCTGCTCA CAAAAAAAAA GTACATCAGA GAGCAACGCG AGRAAGGATT CTGTCATGTC GGCCAATGGG AAGCCAGATC CAAGCTCCCC AGCCTGCCCC TGGTGCAGGG GGAGCTTGTA GGAGGCCTCA CCTGCCTCAC GCTGCTAGGT GGCTGCATTG ATGACACCAT CCTCAGCAGG CAGGGCTTTA TCAACTACTC TOCOGRADACA CARGATECTO ATGRAGOTET TECOCLARCEA GOTECTORARA GEOCETTECTO AGCCGICTOC CAGAATOTOG CICTOAGTOC AGAGGACAAG CITCITATIG AGGCCTCATC AGGCTTCTCC GCCGGGAGAT AGCAGCAGTT TTCCAGGACA алалалала а ACCIGGIGIC COTGIACAAG GATGGAIGTG INCNOTGGCI CCTIGGGAAC TGAGACATAT AGCCCAGACC CACTCCCTGC TCCAGCACCA GCCCCTCCAG CTGACCACCC TGTTGGACCA AGTGAAGAGC CCAAGGTCAA GGAGATGGTA CGGATCTTAA GGGACTGTGC CATTCCTGCC INFORMATION FOR SEQ ID NO: Ε (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45: SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1569 base pairs

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1391

1200 1140 1080 1020

960 900 840

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CGACACCAGC ACCGAATGAT AGGAGGAGAT

720 660 600 540 480 420 360 300 240 180 120

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	269		. 270
			COCAAANTAT ACTATOCOCTT ATGTGAAGGT ATGTGACAAC GTTGACOTCA CCAAATGAGT
	CTGACACTOT TCCGGACTCG TAGCCAGCCT GTTTAGCCAG CCCTGCCCAT AAATACACTC	840	THAACAICA GCICTITITIN CATAIGAAAG CACAITACCT GCICCCCAIT CAAGTAIGTC
V	TOCGTTATTG GCTGTGCTCT CCTCAATGGG ACATGTGGAA GAACTTGGGG TCGGGGAGTG	900	TICCATITATE AGREAGETS ACCACETICA GCAGGAGTEC TECAAGAGTS ECCAACTECE
ר	TOTITICICAC TICOTITICA CIACIDATGA TATTGICAGG TATAGGGCCA CTTGGAGATG	096	CITCCCACAG TACACAAGGC TGTAGTTGTT GTCCTGCAAT CCTTTGTATT TACCTCATTC
	CAGAGGATTC CATTICAGAT GTCAGTCACC GGCTTCGTCC TTAGTTTTCC CAACTTGGGA	1020	TITCCCATCT ANGICCICAC TGAGTTTTAN AGTTAGGGCT GGANAAGCTA TGCCTTACTG
01	COTGATAGGA GCAAAGTCTC TCCATTCTCC AGGTCCCAAGG CAGAGATCCT GAAAAGATAG	1080	GARCAGCAAG GAACCAATTT TTTTCTGAGG GAGAAGACAT TCACCTTCAC TATATGCCTG
	GOCIATION COCHECENCE TROSPEACTS COPPRISONS CACGOOCHEC TRAGGECEACE	1140	GCAGGGCCAC AGTECACAAA ACAAAGATCA GCCTTCATTC AAGTTCCAGG TTTTTTCTTCC
ž	CCCTTGGGGC ACAACCTGCC ACTGCCACAG TAGCTCAACC AAGCAGTTGT GCTGAGAATG	1200 15	TOCCTGAATG ATTACTOCAA AGGGTATATG AAGTAAGAGT TCCCTGTTGC ACATGTACCA
2	GCACCTIGGTIG AGAGCCTIGCT GTGTGCCAGG CTTTGTGCTIG AGTGCTGTTA CATGTATTAG	1260	TCCATAAGGG ATACTATATC GTTTTGCATT CTTCCCCCCA TTCTCCACAT TGTCCTATCT
	TTCCTTTACT GCTGACCACA TTGTACCCAT TTCACAGAGA AGGAGCAGAG AAATTAAGTG	1320	TANGTICOANG COCTITITICAC TCTCAAAAAA AAAAAAAAA TAITITITITIC AGCACTGGTG
70	GCTTGCTCAA GGTCATGCAG TTAGTAAGTG GCAGAACAGG GACTTGAAGC AAGCCCTCTG	1380 20	ITCAAAAGCA ACOTITITAT GOTTAATGGT ITACCAGCAA CIGITGAGAT ITCCAGTIGA
	CTCTGAAGAC CGCGTCCTGA ATTTCTTCAC TAGAGCTTCC TCATCAGGTT ACCCAGAAGT	1440	CTCTTAAAAA TTGCCAATCA TTATCTAGCA GCAATGACAG ATGATTAGGA GCAGTCAAAT
25	GGSTCCCATC CACCATCCAG GTGTGCTTGG ATGTTAGTTC TCCACCCTCG AGGTGTACGC	1500 25	CCICTGAAIT CITICCCIAA TAGGCAGCCA ITIGAGAACT GCACTAGCTG ACATCACTAA
3	TGTGAAAAGT TTGGGAGCAC TGCTTTATAA TAAAATGAAA TATATTCTAA AAAAAAAAA		AACATTATCA GCTAAAGCCA AAACCAAATA AAGGCCCAGA CCAACATCCT GGCTCTCTAA
	алалала	1569	AACCTGTCCA AAATCATTAA GTQAAAGGCA GTAAATGCAG GACTGTGGAT CATGTCACTG
30		30	CHICHGHCHA TGATTARCAA THOGHGACAT GCAACCCCCA TTAAGGTTAA AAGTCCAAAA
	(2) INFORMATION FOR SEQ ID NO: 46:		CHAGICACAC GCATCICITI ATTGGGGAAA AGTGAGACTA TIATGCATTC TTGGTAGGTT
35	(1) SEQUENCE CHARACTERISTICS:	35	TGCAACCTTG CATGAAGAGC ACCCATTGCA TTTCTTTCAT CTTTCAGAAA GCACCGGTAT
			CTGTTCCAAG GGCCTAACAG TACGAAAATA CATTCTGGCA TCACACCTCT GAACCCAAGA
!	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	6	CTGTTCTCAT TAAAAATAAT TTTGGTTTGT AACAAAATTA TGAAATACAA TGCAAGCACC
40	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:	2	TOGGIATAGC ATTATTACTG AAACCACTTA ATTCCCAGCT TTTTGAGTTT TTTAAAAAA
	GGGCCCCCC MCGMKTTTTT TTTTTTTTT TTTAATTAGG ATAATGCCTT TATTAACGAG	09	CCCACTGCAC TAAGATTCAC AATTCATTGC TACATACAAA TTAAAGCTAG TAAGAACACA
45	AARGAAAGGT TCATTCCTCC TTCCACTCCT TCTCGTTGGT TTTCTGGACA CAGCTCACCT	120 45	CTAAGGTCAC AAGTTTCTCA TTCTAAAGTG CAAAAGCCTA ATCATCTGAA AGTGAACAGG
	GATECTISCTA GAAAGSTIST CASTCISCIT STSSCTICCC TECTITAATIS ACTCACGCTS	180	GTAA
ç	TGTGATGTCT TGAGAAGTAT CTATCCACTT CATGTGAATG AGCACTCCAA TATCAGCCAA	240	
₹	CATCAATCAT TCTTACCTAA AGAATAATAA GAANAAGTTA ATATAAAAGA CAAGGGTATA	300	(2) INFORMATION FOR SEQ ID NO: 47:
	AAATAAAGGT TTGAAAATGC TAGTCAAGTT CAAAATTTAA AGAGTAAAAA TCCAGAGATA	360	(i) SEQUENCE CHARACTERISTICS:
55	aagattoggg gtaagttaca gcataaaaa ataggaagaa acttcatggt gogggggaaa	420 55	(A) LENGTH: 475 base pairs (B) TYPE: nucleic acid
	TCTAAAATTA TTCTTACATA AAATAAGTAG ACACCTGAAT TAGAATGAAA ACTGTATTTT	480	
9	CITTIANAITO TAAAAGGCTG ACTCTCAGTT TCACCAGTCT GAGCACAAGT TTGACTGCAA	540 60	(xi) sequence description: seq id no: 47:

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GINGANGCAG CAGGIGATCI TAACTCCTTT CAAAGAGCAG GCCTGTCTGG GAAGCCATGT	THOTILACION CONCONCOC COCCCONTAT GENERATITIAC INSGENTIGAT RETOROCOROC	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:	(A) LENGTH: 1166 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear	(2) INFORMATION FOR SEQ ID NO: 49:		GCCLAGACHA CYTCTGTGAR ANTCTGCTTC CCTCCACAGC TGACCC	GCCCACATGA AGACCAAAGC CAGGACCAAG CCCCMASCCT GCTWAACACG GCAGARTCTT	CCHOCTHANG ATTHACOTYT GACCHTCHAC OTAGGACACT OTGCHGATGG CTACTTGCTG	GAGGAAAAGG ACTAATCAGA GGAGCCAATG AAGTCACTCC ATGAGTTTCC TGAACCCTGC	ATOTOTOMAA TICIOMOCIT ACTOTICIOC CIGOTOGGAC CIGOTOGGA TIGAGAGAAGG	AAGGGACAGA GACCTGGATT CAGATCTCAT TITACAATGA AGACCCCAAT GCAGAAAGTC	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 48:	(D) TOPOLOGY: linear	(B) TYPE: nucleic acid (C) STRANDECNESS: double	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 346 base pairs	(2) INFORMATION FOR SEQ ID NO: 48:			TAAGTAAACT CCCTTTGGAA CTACACAGGT ATGTCTCTCC TTCAACATGT GTGAA	THANGTINTY CHACTAGITG CANGITHAGT GTITCIGITT GTICTGCTTT CCIGTINGCA	TARGUAGUAC AGTIGITATTIC ATTTICTIMA ATTICCTATIGI AGAAGGCTCA GTOTTAGAAA	тослалское остосослся сясьлюсого сясясясяля стосемсяс алатосясял	AGAACTOTOG TOCCAAATTO CTOOTOTCTC TCTTTGGAGA AACCAACCAG ATACATCTOC	OCCCAGTAAG AACTOCTOTC ATGAAGGAGG GGCCACCTTG TAAGAGACAT CATTACTACC	АОЗЛАЛОССА ТТОССАМТУС ТОЛОСССТТО АЛОООСЛАЮ ДООЗЛАЛСАО ТОТТАССЛОЛ	тостогозос сселдаламс амереалскае теалалисьме оскленеает тетитескее
180	5 6					346	300	240	180	120	60								475	420	360	300	240	180	120	60
60		55	50		45		40 ,		. <b>3</b>			30 ,	•			74	20 c	13			0	10 1			.s.	0
сититилос таатадыте ситетьова ессавесоге тетпоиства твототтесе	AUGMITTITA COTCTOTTAC AMMOMBATT TEOCHGAING CECTIVISOGI CCCACTOTIC	GCAGTAATTC CTGTTAGCCA CTGCATCCAC CAAAACTAGT TTATTTTTTCC CCTCAAATTC	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1405 base pairs	(2) INFORMATION FOR SEQ ID NO: 50:		ATAMACCCTA TITCITATIT ATAMAMAMA AMAMAMAMA AMAMA	AGCTOTOMAC AGCAGGGGGT TOTOTOTOTOTO TICTOTITICT CIGCTIGGCG AACTITICICA	TOUTHINGAAA GCACAGAGCC TEAGGGGGGCC TGGGCCACAGA ACACAACCAT CTTAGGCCTG	GTCAGCCAGA GTTCTGAAGG CCATGCTTTC AGTTTCCCTT GTTGACAATT GCTCTCCAGT	ATTITICICIC CTOSTAGCIS AACAAAGSIC TAAATTAGCI TAACAAAAGA ACAGGCIGCC	AGGINAAGCA GEATEANIGGT GITTINAGAC CAGAGCITIGG GACCAGGGCT CCTACACCTA	CCCGCCTTGG ACGTGTTTCT TTAACCTCAT CCATATAATA GGGCCGTGGG ATGGTTGTAG	TTTOTCAGGT TOTCCTTGTT TOGATCCCTC AACTAGGTGA TAAGCACTGG AGGGGGATGA	GIGAGICTGA CCRYGAGSCG GRCCCCTICA CCTIGGCIGG GCIGGICCIG GICCTIAGGI	ACAAAGTTCA CGTRGCAGGT CTRGGCAAAG ACTGGGCAAT TGAGCAGAGG AGACGGACCT	GOCCTIACTT TICCICCCAC AAAGGAGICG CAGCCACGCT AGCICIGACT IGCCACIGIG	TOCTOTOCTIG COTTOCCTOG CAGTOTTCTG GGGGTGGATT CCCTACANCT AGATGTTCAA	TTOTOTTOCT GTOCCHOOGG ACCOTOGAGA ANOTOTOLAG GOCCOCTCAC TOCAGCAGCC	THEATROOCEA GOSCIAAGINE GOSCIAGOTEA GCCAGOTECT GCCAGCOCINE TETETEOGRAC	CTCACCTAIG COCAATGANA GITATTGAAG GACTGOITGT TGATGITGGT GAGCGTAICC	TOOTTOGOCCA GOCCCCTOCA TOOGLAAGGGA GCCTOCTOCG GOOCLAGOCCA GCTTOGOGOGTO	AGTOCOTUAA COTOCCACO COAMTIOCOT CAGTITOTOCT GAGCOTCATO TOTOCCOG	TITUSTITAG TEATETAGAG TEOTETOGAE TAAAGGTETT TEAGGTETEC TIGGESTOTG	GOCCAAGOTT CTATTOTAAC AGTAGOCACA GTATAGTOGG AFCATCACAT CAGCTGOOTT	OCTORACIAGO CACAGORAGO COTOTOGARA TOGRITCADAR ACTOROTOT CAGODRAGORA

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\$	Jd average vin	PCT/11/S08/f04493	WO	WO 98/39448 PCT	PCT/US98/04493	14493
•	273			274		
	TITIAACCCIT GGCAIGTATA ATAGAATITT GGTGAATGAA AGAACCCAAA TAGGCCAGAT	240	F	TCACCTAGGA ACATATCTCT GCGGTCTCTC CTGCTCTCAT AATGAAGACA TAGCCGATTC	240	
	AGICCCCCCA GGCCCTGATA TCCATAAAAG GCTTGGGAAT GCATTATGTA ATTGTCCTTA	300	F	TOTOCCOGGO CCCCTTGCTG ATGCTCCTCC GGGTCTGCCT CGGGCGTGGG TCTCTGGGGA	300	
ς.	STCTTTTTGT TSTTTTMGAN AAAAAAACA AGATGGGCTC AGATGGATGC CTAGGTAAAA	360	ر د	CCCTCCAGAG GTGGAGGTGG GCTGATTGCTC GTGGTTGATG GTTTTTGCTCC	360	
	ATGCTTCCTA GCTGTGTACT CATAACTTTT CTTTGAATTG AGTAGTGAAA GGAAGGAGGA	420	ō	CCCTACCTTT TTTTTTGAG TTTATTCTGA TTGARTTTTT TTCTTGGTTT CTGGARAAAC	420	•
9	GGAAAGGAAA TTAAATGTCC TTCTAGTATT CTCTGGACTC AAGTCTGACA TATGAGATAA	480	01	CACCCTCTOS GOACAGARA ATAAAACATG TAARATTTTT AAGAAGGAAA AAAAAAAAA	480	
2	TAACCTATAT TGAAATGCCA AGAATTGTAT CTGAAACAAG AGAACAGTTT GACACATTTA	540		ABABARMIT CONTOUNT FORK	3	
	TCATCCCTTC ATATTACATA TTAACTGAAA CCAAFTAATA AACATATGAA ATATCCATTG	009	ŧ	AND THE CONCESSION OF THE CONC	on on	
15	CACAAGGCAA AGGCACTAA ACCTITITGTT TCTTTTTTCTA CATAGCAGAA ATTGATTTTT	999	15			
	TITITIAITIT TITAGGGGA CCTATATAAT TATGACCCAG TGATGTCTTT TGGTGACTTA	720	Ç	(2) information for SEQ id no: 52:		
;	AGCTIATGAA TTCAGOTIAC AATTGAGTTO ATTCTAGATO GTTACTACCT TGAAAAGGAT	780	20	(1) SEQUENCE CHARACTERISTICS; (1) IDMITH, 777 home main		
3	STITICOTICCT TATIOTISACAC GACCAGAGC CTOCTICOGGA ATAMACAAAG CAGGITTICAL	840	3			
	GCCAACACCA ACTOSTAGCT TTAGTSGGCA GATGGGGAGT GOTTCACAGA CTTCCCAAAA	006		(D) TOPOLOGY: linear		
25	TOTOGOGGET TTGGGATTTT CCACACATC CCACGTGTGT TGTTCATTCT TCCTCTTTTC	960	25	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:		
	ACACTETTOS ATGGATMATT TGRAAATOOT GRAAMMINGY YYKRAATTTG CCCAATAGCC	1020	2	naagtatett goccagtta ttacagagga cgataaatga ttecatgtgg atagggcata	9	
ç	WIGHGCCACC ATTCTTWATG ACACCATAAC CAAATAGTTC CWIAATGTTG AAATATTACA	1080	λ Σ	ACATACAGAG AATGAGACTA TGCCAGAAAT GGGAGGAGGC ATTTGAAACA ACATGAGTAT	120	
8	AACCIGITAC CAGOCHSHA KIWACCOMA WITTICCCAT GITIGIGAA ITGATAITGA	1140	_	CTCAGGGACA GATGGATTGA TTCTGCTATT GGTAGGCCTG GAAGCAANGG TCAGAAGTAG	180	
	ANTAGCAGG CTAAGGAATT ACTGGCAAGT TITTAGCCTOT GOGTAATACC TTAGGGTTAT	1200	ð	CHANANTIGG ATACCHANAG CACTATINGT CACCCHAGCT ANGTGGIATA GCTGGCCCAG	240	
35	TTAAATATTT GTAATTTTAT TTAAANGTTC ATGAATGTTT GAAAGGAACA AAATTATCAG	1260	35 T	TAGGAGAAAT GCAGGTTTTG CTCTACACTA AGTTCTCCAA CTCTTGATAA GCCTCCAAAA	300	
	GGATGGCTCT TIGCCATGGG TCTTATTIC ACCCTCTTT CIGTAGGAA AAAGAACAAT	1320	¥	acaaatetta goggaaaaaa acgcagctgg ttatgaaaag atatatetca tttecattaaa	360	
:	GICTIAATGI ATITITIAAAG TITITIGGIAT AGITICIAAT TCCAATITIA ATAAAAGITI	1380	. ₹	ANATCAATGT CAATGCTGTT AATAGAATCC TTTTATCTTC AGGACAGAGG CAATGCCCTA	420	
9	TWIRITAADAA AAAAAAAA AAAAA	1405		AACAAACACC AGCTCAAGAG CCTCTGATGC CAACCTAGAG GGTACCCAAA CACAAACTTA	480	
			8	GCATAGAGGT AAGAATCTCT ATGTCTTTTG GTGGAGGCAA AGCCATTTGG TTGGTACTTC	540	
45		4	45 AC	ACAGGAACAT CITTICTACCA AGTCITICATE ATATGGTATE TGCCAGGAGT CICCAGTIGT	009	
-	(2) INFORMATION FOR SEQ ID NO: 51:		Ē	TTGCACCACT GTGTCATAGC TGAGAATACG CTGAAAGGTT AGTTTTGATC CTGGAAACCT	099	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 504 base pairs	v	ž C	ATTIACAATT GCCAGCTGAT GTCCCTGCTG CCACTTAAAA AAGACTTGGG TCTGGCATAG	720	
S	(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: Linear	•		GCAGAMAGGC CITOTGGTCCC CITOTTGCCCGA TICINOGCTC GAGGCCAAIT NCCTIAT	יור	
	(x) spousoce description: sec id NO: 51:		į			
55	ACCOMPANY AND AND AND AND AND AND AND AND AND AND	C	ე ლ	(2) INFORMATION FOR SEQ ID NO: 53:		-
	TOTAL TOTAL	3		() CENTED THABACTED CONT.		
		120	ξ.	(1) SEQUENCE CHANNELEKISILGS: (A) LENGTH: 602 base pairs		
9	AACTITICAGA GAATTACTAT TTACTITATT AACTTACGGA TTTATTATAT AAATATATAT	180 6	<b>3</b>	(B) TYPE: nucleic acid		

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CARGOTTICG GARGAGCAGG TCAATAGITT TGAGAGGGAG TITGITCCIT TITTITITCT CITAAAAATA CQAAAATAAA TITTACAAGG TICCGGTITT GGTGGTGGAA AGAGTAAATT CATTITICGAA TACTOTICGAA GITTITATCIC TIGCATATAC TITATACIGA AGTATITACIC AAAAAAGAAA AATTAACTGT AGCGCTTCAT TATACTATTA TATTATTATT ATTATTGTGA TICTCACGIA TAITIACCIG IGACIIGIAT TIGITATITA AACAGGAAAA AAAACATICA GCCAGTTTTT TOTTACAGIG ACAGTATOOT TACCIGCCAT TIAATATTAG COTCGTATTT AACCTIGGAT ACCCCTTTIC TATCAAGIGC TITAAAATAT AGACTAAATA CACACATCCT CATTATACTC TTAAATTGTT GTCAGTTATC AAACAAACAA ACAGAAAAAT TGTTTGGAAA ATGACTACAG TGTTATACCC TCCAATCTTT GCAGGTGGGC ATGGAACACT GCTTGTATCA GATTATICCC AGGACAGCIT CAGCAAACAC TACAAGICCA CGGIGGGAGI GGATITIGCI CCCGACCACA CTTCCCGCCT CCCTAAAACG CACACCCCGC TAGCCATGGG CAGCCGCGAC CCTCACTTCA GTTTGAAGAG GGTCCGGATC CAAAGGGGTT AAAACGAGCG AACCCCGATC 2 ATTATOTTTO ACOTTACCAA TOCCACTACC TTCAGCAACA GCCAGAGGTG GAAACAGGAC CTGAAGGTTC TCCAGTGGTC TGACTACGAG ATAGTGCGGC TTCAGCTGTG GGATATTGCA CACCTOTTCA AAGTOCTOOT GGTOGGGGAAC GCCGCAGTOG GCAAGACGTC GCTOGTOCAG AGTCACTGAC TIGGAGCCGC ICGGGGGAAG ICCCGCCCAG ACAGGCGGIG GGIGGGAAIG GOOCAGGAGE GETTEACETE TATGACACGA TIGITATIATE GOGATGEETE TGEETGTGTT CTAGACAGCA AGCTCACACT ACCCAATGGA GAGCCGGTGC CCTGCCTGCT CTTGGCCAAC AAGTOTGATC TGTCCCCTTG GGCAGTGAGC CGGGACCAGA TTGACCGGTT CAGTAAAGAG INFORMATION FOR SEQ ID NO: 54: Ě (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 53 (A) LENGTH: 1749 base pairs
(B) TYPE: nucleic acid
(C) STRANDEENESS: double
(D) TOPOLOGY: linear 9 9 TOPOLOGY: linear STRANDEDNESS: double 7 600 540 420 360 300 180 480 180 120 600 480 420 360 300 240 60 6 330 23 20 55 50 25 ᅜ 5 8 GIATITITIGA AACITICIAAC GICAIAATIA AGITICICIT GICITGGCAT CAAGAAIAGI TICICATCAG CCCICAATIT GIGAICCGGA ATTITIGIGAG AAGGATIAGA AAICAGCACC GOGCCCATTO TCACTINGAA ANGACACCTO GAACCCATOT GCATTTCTGC ATCTCCTGGA CCCACCIGAC CATITIATIA AGIACATITG AATIGICICC TGACIACIGI CCAGIAAGGA ACCCAAGGG ACTACATCAA TCTACAAACC AAGTCCTCCA GCTGGTCCTG CTGCTAGTAG AACGGTTTCA CAGGTTGGAC AGAAACATCA GTCAAGGAGA ACAAAAATAT TAATGAGGCT CGAGATCATG CCACCGCACT TCAGCCTGGG TGACAGAGAA GGACTCCGTC TCAAAAAAAAA GGCAGGCGGA TCACATGAGG CCAGGAAITTC GAGACCAACC TGGTCAGCAT GGCAAAACCC CAAGITITIT GOCCGOOCAT GOTGCCTCAT GCCKGTAATC CCAGCACTTG GGGAGGCCAA TIMOCCITIC ACAIGIIGCI GRCICACATI AGIGCCAGII AGIGCCIICG GIGIAAGAIC IGITIGACIT ATTTICCATC CCAGTICIGG GAGGICTITI AAGICICTIC CCITIGGIIG ATGAGAGICC TCATTGAAAA GATGATGAGA AATTCCACAG AAGATATCAT GTCTTTYTCC GOGTOGCCTG TOGTATIATOG AAAAGTAGCA GGGTGGTCAG GGTGGGAGAC ACAAGATGTT CCICCIGICI ACIGGCICCA AATAGACCAI GICAGCIICA CCCCCIGGCI TIGIGICIAI CATTANATTT CITACAGIGA ACIACATATI GICCATAAGI GCTICATCAG GACTCATCGC AAAGAGATIGG GCTCTTTATT TICTCGAAAA ACCAATTIGG AGTTACTCAT TITTTCCATAA (2) INFORMATION FOR SEQ ID NO: ACAATCNAA AAAAAAAAAA AAAACTCGAG GGGGGCCCG GTACCCAAAT CGCCSTGATA GTGATCGTAW TACTOTOGAG ACTUAGOTOG GAGAATOGOT TGAGACTOGG AGOCAGAGOT TGCAGTGAAC COTCTCTACT AAAAGTACAA AAATTAGCCA GGCGTGATGG CACGTGTCTG TAATCCCAGC AGAAGGAGGG GTCTGGGCAT CTGTGGATTT TTGGCTACTA GAAGTGTCCC AGAAGTCACT IGIAGOCCOT IGITICAGAT ICTTICIGIC TIGGAAIGTA AACATCIGAT ICIGGAAIGC TIGATATCAT GACTICCAAT IGAGAGGAAA AIGAGAICAA AIGICAITITC CCAAATITICT IGCGTTTTAG AGATCATAAT TCTCACCTAC TTCTGAGCTT ATTTTTCCAT TTGATATTCA ε (xi) SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear (B) TYPE: nucleic acid
(C) STRANDEDNESS: double Ξ LENGTH: 1896 base pairs 276 55 1740 1620 1560 1500 1440 1380 1320 1260 1200 1680 1020 960 900 840 780 660 240 180 120 69

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(2) INFORMATION FOR SEQ ID NO: 56:

25 33 8 5 8 S 으 2 8 8 360 38 420 480 240 9 9 22 380 840 1740 1800 1860 8 98 1020 1080 1140 1200 1260 1320 1380 1440 1500 1560 1620 1680 TITIATAGICT AGAGCCITTIA AAAAACCCAG CAGAAIGIAA TICAGIATITI GITTIATIIGGC TOTITITIES CAGATTOTIC AAATTAAATG AATTGAAAGG GAAACTCAGA GTACTAGGAC TIGITICCIT GAICTOICAG AGGICACAGF AACCIGGGC GAGCIGITAT TAITTAITAT GITTATTAAA AGGAAAAAA TOTCITGCAA TGTGCTGTAA TCACAAGAGG AGAAAATAAC ATAATAGTAG TAGGAAGTTA ATAACTGGTT CYCTGTGTTC CAAGCACAAT ATTACAACTT CITITIGAACC GIAAAIAICA GAAIGAAICC ICITICCCAGG GGAITIGAACA GAAGCITIAAI GITTACAAGI GITITGAATITI GIGATCIGAA ATAACACAAA AITAAAAACA IGAITITCICI AATTITICCAA CTAGAGGAAG AGAAACTIGI GGAAAAGIIC ITITITITIC ITITITITIT CTTAAAGAAG GGCAGCCAAG GTAGTAACCT AAAATAGTG CCCAGGCATA TGAGAGTTGT CCTACGAGGT TAAAGAACAC ACTGTTCCAC TGTATGCCTT TGCCCTGAG TGGCCAGGGA GOTCAACTIG ACCCIGCCAT GITGOTTIGA CITACIAAGA CACAGGAAIC ATTOTITICC ITGACCAGGG TCTCACACCC TGGAGGAATG TTAAGTAAGA GAAAGAACCT CTTTCCTGAA TATTCACATG TAAAAGACCA AAGTAATTTT TCTGAACTTC TGCAATTCTG AGAACTCTCC AAGGAATTTA CAGTGATTTT AGTGCTTGTC AGCATTTTTC CATGAGGACT TTCATACATT IGACTETITIA GITCACAGGI TCCCAITGAT IGICAGCAAG ATAITITAICI CIVITAGCCCT TOGGGATCCA GCTCAGAGCA ATCTCTTGCA TTTTTTTACC COTGTATGTA CAGATATCAT INCITIGITIA TECCATGACT TGAAAAGIT TGGGAAGCTC TITAGCAATA TCAGCTAAAA AAAGGAAATA TAGTCCTAAT CTTTACTATC CACTTCTAAA TTTTAATGTGA ATTTCATACA IGITAITAGT IGITITICITT ATAATITIAT AAAAATIAIT CAICGGGAGT ITAACTICCA ATCCATTTGC ATAATGACCC AGATCATCAT TITICTGCAAC TGAGAATTAT ATTITICATCAT IGCTICTAGA AGICTGCAAT ICTITACTITI TCTITIGGIGC ATTAITAICT AGGIGCCAIC GAAACCCCGT CTTCTTACTT AAAATACCC AAAATTAGGC CAGGGGCTG GATGGGTGGG GIGCCIGITIA ATCITICAGCI ACITINGGGGA GGGCTIGAAG CCAGGGAGGA ACITCCCCTGG GGATATGAAA TCACAGGTGA TAGCAGTTGT CATTCAGTAA TTTCCTACAA GCAGCACCC CTTCCATGCT ATCGGATGTG TTGGGCTCCA TGCAAGAACT TGGAAGAAAA ACAGGCAGGA ACTOGATAAT GTGGAGTGAC TAGAGAAGTC AYATATCACT GTAAGGTACA GTTAGGGGTA ACACTITIAGA GGITTIATIAT TITITAAAAA CITITICITIGA ACTOCTIGGGC CAACATGGGT

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8 120 180 240 300 360 420 480 540 9 99 720 780 840 900 960 070 98 140 200 260 320

CYATACAACT IGITITIAACC ITICAGAACA ITICACAGAAA ITAIGCAAIG GITIGITIGAG AIACGGACITI GATIGOTICOTO TITAATICAGT TITOCITICCAA AGTIGOCITAC TICAAGAGGCC CTAAGACTIGG IAGAAATTAA AAGGATTICA AAAACTITICI AITICCITTICI TAAACCTACC AGCAAACTAG GALTICIGALA GCAATGAATG GIATGAIGAA GAAAGITITGA CCAAAITITGT TITITIGITG TATTCCCTTT AGAATGTTT ATGTATGAGT CACCTGTTAC AGAGITICAG ATCGGICACT GATAGIATGI ATTICTITAG TAAGAATGIG TIAAAATTAC ICTITITIAAA ATAGACATITI GIGGGGCTCA CACAATAITAT GAAATAGTAC CCTCTAAAAA agagaaaaa aaaatcaggc ggtcaaactt agagcaacat tgtcttatta aagcatagtt TATTITCACTA GAAAAATTI AATATCAAGG ACTATTACAT ACTICATTAC TAGGAAGTIC ITITIVAAAT GACACITAAA ACAATCACTG AAAACITGAT CCACATCACA CCCTGITTAT ITICCTIANA CATCITIGGAA GCCIANOCTI CIGAGAATCA IGIGGCAAGT GIGATGGGA STANANTACC AGAGAAGATG TITAGTAGCA AFFANAGGCT GTTTGCACCT TTANGGACCA SCHOOCCHOT ACTGATTCCT GOOCCCAGAG TGGCATTATG ITTITACAAA ATAATGACAT ATAGAAAGTA TTACTTTICTT TCATATGGTT TITGGTTCAC TGGCTTAAGA GGTTTCTCAG AATATCTATG GCCACAGCAG CATACCAGTT TCCATCCTAA TAGGAATGAA ATTAATTTTG IAICTACTGA TAACAGAATC TGGGTCACAT GAAAAAAAT CATTITIATCC GTCTTTTAAG INTATETITA AAATAATAAT ITATETEET GCATAITIGCA GAACAGCTCT GAGAGCAACA GITITCCCATT AACTCTTTCT GACCAATAGT GCTGGCACCG TTGCTTCCTC TTTGGGAAGA GGAAAGGGTG TGTGAACATG GCTAACAATC TTCAAATACC CAAATTGTGA TAGCATAAAT AAAGIAITIA IIITAIGCCI CAGIAIAITA ITAITITAAIT IITTAGGIAA IGCCIAICIC TTGGTCTATT AAGGAAAGAA GCAATCAGTA GAGAATTCAG GATAGTTTTG TTTAAATTCT ATGREACATG TITIGEATGIT TGITITGETTG TIGAATTITIT GAACAGCEAG TIGACCAATC IGCAGATTAC ATGITTITIAC AGTGGCCTGC TAITGAGGAA AGGTAITCTT GTGAAGATGC TAGGGAAGCT ATGCTCAGAT ATTCATCGTA AGTCTCCCTT SEQUENCE DESCRIPTION: SEQ ID NO: SEQUENCE CHARACTERISTICS: (A) LENGTH: 1753 base pairs (A) LENGTH: 1753 base pair (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear PROFIGINGE INTRAATING AAATCAINCE Œ, 3 25

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1380 440 1500

AAIGAICITI TAAAAAGAIG AIGCAGIICI GIAITITAITIG IGCIGIGICI GGICCIAAGI

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ANCCCCGGGG NGGCCCAGNA GGTTTGCCAG TTGAGT

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GGAGCCAAIT AAACAAGITT CATATOTATT TITICCAGIGI IGAAICICAC ACACIGIACI

1560

TUGGTTTTAA ATUGACATTU TCTGTACCAG CTTCATTAAA ATAAACAATA TTTGTAAAAA

1200

CATGIGATTA CITCACICCI GGACIGIGAC ITTICAGIGG AGAIGGAAGI TITTICAGAGA TICAACAGIT TAGATATICT TITTATTITT TITCITTICC CICAATCCIT TITTATTITT ANTGAATTICA CTCAGGITTIC TCTTTGAGGG TCAGAGAATT GCTGATAATC ATACTCCAAA GAAAATGACA ACACATCTCA AGAAACTCAA AGAATCATAC TOTCAAAGAC AGGGTGTTCC AGGIGAATAT ATTAAACTCA AAGICATIGG ACAGGATAGC AGIGAGATIC ACTICAAAGI COTCATCATO TOTGACCAGG AGGCAAAACC TTCAACTGAG GACTTGGGGG ATAAGAAGGA COGAGOCGAGG TICTOCTIAC CCGAGGCCGC TGCTGTGCGG AGACCCCCGG GTGAAGCCAC GCGGAAGTTA CTGCAGCCGC GGTGTTYGTGC TGTYGGGGAAG GGAGAAGGAT TTGTAAACCC TCCTTGAGAT TTCACTACCT TTATCTTAAA AGTTGTGTAT AATTGTTAAA ATCTGTGAAA TIGAAAATIT CCTICCAICC IGAATAACGA ATAGAAGAGG CCATATATAT IGCCICCITA ANATANGATO GACCANTICA AGTOTTCATA ATGACTITICO ANTIGGOCCI GATGITICINO TTACGIGIGC ACAGAGAGGT CACCITITIC AGGACATIGC ATTITICAGGC TIGIGGIGAT TTIGAAACAT CIGGIAATIT GAATICIAGI GCICATIATI CATIATIGIT IGITTICATI agaactogga atggaggaag aagatgtgat tgaagtttat caggaacaaa coosssstca GACTAACTCC AMAGATOGCT TCACTGAAGA AAAGGCATTT TAAGATTTTT TAAAAAATCTT TOGACCAAAA GAAGAGGAAT ATCAGGTTGA AGTCAAGATG ACAGATAAGG TGAGAGTAAT ACTGAACTOT GGAAAAATGA CCTTTCCTTA ACTTGAAGCT ACTTTTAAAA TTTGAGGGTC CTITIGIAAT GIGGIGITCA AAACGGAATT GAAAACIGGC ACCCCATCTC TIGGIGATICA AGCCICAGIC CCCTICATAT TACCCICICC TITITIAAAAA 1753 1740 1680 1620 1020 960 900 780 720 660 600 540 480 420 300 120 6 3 8 35 30 25 2 5 5 55 8 TOTTICOTIGG CICCICCIGG TIGGCCITGT CIACCIGGIG TCCCACCIGA GICAGCGGIG CANCENTECT GTOGRAGEEGG TEACCTECAT CETGETECTC TTECTGETCA TEATGETTOG CCCCTTCAAT GACCTCAACC GGCAGCTGGT GAACATGGGC TITCCGCAGT CCAGCCACCA AGAGGTAGCA TICCTCGACA GAGCTICTIC AATAGGGGCC ATGGTGCTCC тсамалалал алаллалала GACGTCTCCT CTGCCTAGGG AGCAGGACTT GGGCTTAGGG CAGGTGGAAA AAATTCCAGA TTATOCCOAG AAGATOTOAG CTOGATOCOA ACATOTTOCO ATOCCTOTOG AAGACATOCO CATTAAGTOG CACAAAATCA GAGCAAGAAA GCGATGCCCT TCCCAATTCT CTCAATCCTT TICCCAAAGI GIGGGIGGGI CCGTIGGTIC CCGAGATACI TITAGGIGGI AIGGGGCCIG ACCICIDADO GCIDATAGOS GIGOGITITOT IDADADOGAC ITIOCIDOGOCO ITIOGIOTICAG CCCAGGGGT CCTGGCCCCC GCCAGCAGCA GGCAGGTGCC AGGCTGGGTG CTGCTCAGTC GOGGATICTIG GCCAATIGGGG AAATICGIGGCA GGACGACGAC CCCCGAGIGA GGACCACTAC CAAGCCTCCC TCAATTTCTG GTOCAGCCCA TCAGGGACCC ACAGCGCCTG GGAGGATGGT TOGOGOCTIOC AGACACAGOA TOTACTOAGO GTOGOTOACO TOTOTOAACA TOACTGACTG 2 AAAAAAAAAA AAAACTCGAG GGGGGGCCC ACAAGIGGIA GIGGCATICI AGCAGGCATA TTTGGAGGGG ATCTGGTGGT GCCTTGAAGG TATGATCAGA GAGGGGACCA INFORMATION FOR SEQ ID NO: 58: (i) SEQUENCE CHARACTERISTICS: £ ACTOTTTTIG TITICCCCTIT GIGITAAGCG TGAGGCAGAG GGAGACGITA GICCCAGCAT SEQUENCE DESCRIPTION: SEQ ID NO: 58: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1049 base pairs ATTTATTOTO ACCITTICAA TAAATAGATT TAAGTAAAAA TITTAATGGT ATATTITTAT TGGCTACTTT ATTGTTTAGG GCATCTCGG 1220 1049 1020 960 840 780 720 660 600 540 480 420 360 30 240 180 900 60

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(X

SEQUENCE DESCRIPTION: SEQ ID NO:

57

(B) TYPE: nucleic acid(C) STRANDEDNESS: double(D) TOPOLOGY: linear (A) LENGTH: 1220 base pairs 5

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 57:

5

AAAAAAAAAGG GGG

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AAAAATAGTT

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8

AAATGAATAC AACAGAACAC TOCTCTTTTT GATTITIATTT GTACTTTTTG GCCTGGGATA

1140

8

(i) SEQUENCE CHARACTERISTICS

(2) INFORMATION FOR SEQ ID NO: 59:

GTCAGAAGAT CCCAGAAAAG TTCTAATTTT CATTAGCAAT TAATAAAGCT ATACATGCAG

			0770000000	Ĭ.	PC************************************	1077
\$	WO 98/39448 PC 281	PCT/US98/04493	VO 98/39/446	282		
	;					
	(A) LENGTH: 1776 base pairs (B) TYPE: nucleic acid		GGGTATATAG ATTK	GOCTHINING ATTOTATINA AMMAMANG CTHINITATICC ATMINICTHE ATHINATHIG	1680	
		v	ACGCAGAAAT AAA1	ACSCAGAAAT AAATCTATGA GAAATCTATC TACAAMMAA AAAAAAAAA AAAAAAAAA	1740	
S	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:	0	AGGAATTCGA TNTC	AGGAATTCGA THTCAAGCTT ATCGATACCG TCUACC	1776	
	AAAGAGGATG TGAAGCTAGA GGTCCCCGAT GGCTGGTCGG ATGGGAAGCA CAAGGCTGAG	. 09				•
9	GOACTGGATT GTAAAGGCAC TAAGTCGTTC TGGGGTGAGA ATCAGACATG GGGGACCTCT	120 10	(2) INFORMATION	(2) INFORMATION FOR SEQ ID NO: 60:		
	ACCITICACAT COTOTITICOT TOCAGSTOTG GACATOCOGA GOOGLAGTOC COCACACTOA	180	)OBS (1)	SEQUENCE CHARACTERISTICS:		
ž	GTGCAOTGAT GAGTGCGGAA GTGAAGGTGA CAGGGCAGAA CCAGGAGCAA TTTCTGCTCC	240		(A) LENGTH: 443 base pairs (B) TYPE: nucleic acid	•	
3	TAGCCAAGTC GGCCAAGGG GCAGCGTIGG CCACACTCAT CCATCAGGTG CTGGAGGCCC	300		(C) STRANDEDNESS: double (D) TOPOLOGY: linear	<del></del> -	
	CTIGGTOTCTA COTOTTTIGGA GAACTGCTIGG ACATGCCCAA TOTTAGAGAG CTGGCTGAGA	360	(xi) SE	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:		
70	OTGACTITICS CICTACCTIC COGCIOCTICA CAGIGITITICS TIATOGGACA TACOCTICACT	420 20	ACAGATAAAT AAAT	rchgathaat aartaarta taartaaat taartaaaa arctgagcta atctgagtaa	. 8	
	ACTIAGCIGA AGCCCGGAAT CITCCTCCAC TAACAGAGGC TCAGAAGAAT AAGCTITCGAC	480	ATTGAGAGAT TTCA	ATTENDAGAT TTCACATGAA AGCAGGATT TCTGGCTTCC CAGGACAGT CAGAAGAGCT	120	
2,5	ACCICICAGI TGTCACCCTG GCTGCTAAAG TAAAGTGTAF CCCAFATGCA GTGTTGCTGG	540 25	AGCTAGGAAC ACTGGTCTGC	GICTGC TIGGCTACCT TCTTTGGAAC AACATGAAAT CTAGCTCCCT	180	
3	AGCTICTTOC CCTOCCTAAT GTGCGGCAGC TGGAAGACCT TGTGATTGAG GCTGTGTATG	009	PITT TITITITY	TITITITITI ITITIOGOCC ACTICATOCA TICACAIGAC CIGCOTOGOC ICTICCAGGIA	240	
	CTGACGTGCT TCGTGGCTCC CTGGACCAGC GCAACCAGCG GCTGGAGGTT GACTACAGCA	099	AGTGAGTATG CAAC	AGIGACIATO CAACAAAAT GTAGCACAGG TTTTIGTOGGT GAACTAGGTG GTTTCAGGTC	 	
30	TOGGGCGGGA CATCCAGCGC CAGGACCTCA GTGCCATTGC CCGAACCCTK AANAAAACC	720 30	CAGCTCTGCC ACT	CAGCICIGC ACTIGCTAGC ANGACCICOF GCGGAAFTCC NGCAGGAAGF TTTTTTTTT	360	
	ATTANAGITIA CGACGCAGC AGCAGCCGCA GCCACATCTC AGGACCCTGA GCAACACCTG	780	TITITICAGIG CICC	TITITICAGIG CICCAGICCC CCIAITIGGAG AATOCIGCCC CCCCTGGGA CAGAAIGITC	450	
90	ACTERACTER GEGRACCIAC TOCTGGCACC AACCAGGGC ASCCAGCAAG MAGCCTCAA	355	ACCTEGCCC CGCGANTCCC	ANTICC TGA	443	
3	AGGGCAAGGG GCTCCGAGGG ANCGCCAAGA TTTTGGTCCAA GTCGAATTGA AAGRACTGTC	006				
	STITICCIECE TOGGGATISTO GESTICCEAGE TECCTOCETO ECTETTAGA STECTEAGAG	096				
40	AGCTITCTOT GCCCCTGGC ACCTGAFAAT CCTAGGTTCA TGACCCTTCA CCTCCCCTAA	1020 40	(2) INFORMATION	INFORMATION FOR SEQ ID NO: 61:		
	OCCCAAACAT AGATCACACC TTCTCTAGGG AGGAGKCAAA TGTAGGTCAT GTTTTTGTTG	1080	(i) Sequ	SEQUENCE CHARACTERISTICS: (A) LENGTH: 2888 base pairs	·	
7	GTACTITICIG TITITITICIGA CITCANGIGI ICCATIGGIC COCCCIGGOA IGCICICICC	1140 45				
3	CITETITICCI TAAGAGCICA GCATCIOTCC CIGITCAITA CAIGICAITIG AGTAGGIGGS	1200		(D) ropology: linear		
	THACCCTICAT GGGGGTCGCT CTGTCTGGAG CATAACCCAC AGGCGTTTTT TCTGCCACC	1260	(xi) SE	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:		
20	CATCOCTICA TECCTGATCC CCAGTTCCTA TACCCTACCC CTGACCTATT GACCAGCCTC	1320 50	TTAATGTTGT	CAATAACCAC CAGGCCAAAC AGAATTTATA TGACCTGGAT GAAGATGATG	09	
	TRANSPOCCA TAGGGCCCCC ACCITTACTC ACACCCTGAG AATTCTGGGA GCCAGTCTGC	1380	ATGGTATAGC TTCC	ATGGTATAGE TTECGSTTEET ACTANACAGA TGAAGTTTGE AGCCTCAGGE GNETTTETEE	120	
;	CANGCCAGGA GYCACTGGAC ANGINCANCC TAGAATCCTG TCACACTACA GYCANTHCTT	1440	ACCACATGGC TGG	ACCACATIGGC TGGGCTAAGC AGTTCCAAGC TTTCCATGTC CAAGGCCCTC CCTCTCACCA	180	
55	TTCCTCTCTC TGGCCCTTGG GTCCTGGGA TGCTGCTGCT TCACCCCAG AGCCTAAGAA	1500	AAGTGGTTCA GAAT	AAGTGGTTCA GAATGATGCA TACACAGCTC CTGCTCTCCC TTCCTCTATT CGAACAAAG	240	•
	TOGCHOCCGT TICTTAACAT GTTGHGAGAT GATTCTTTCT TGGCCCTGGC CATCTCGGGA	1560	CCTTGACCAA CATC	CCTTGACCAA CATGTCCCGG ACACTGGTGA ACAAGGAAGA ACCCCCCAAA GAGCTGCCAG	300	
99	I ACCITICATIGG CAATCCTIGGA AGGGITTAAT CTCCTTTTTGF CAGTITGGTG GGGAAGGGAA	1620 60		CIGCIGAGOC TOTICTICAGO COATIGGAAG GCACCAAGAT GACTOTGAAT AATOTGCACO	360	

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GOCTICTIGAGG CTGCCCAGGA CGGGAAAGTC CAAGGAAGGG GCCTGGTGGT GCTCCACTTG

TOCAGOGTO AGTOTOTOAG GCAGGAAGGA COCAGGATTT GAACCCAGOT TOAGTGTGCA

CAGTICITIA AAGAAIGCIG CITITIATIC ICCTAACCCI TICAAGIGGG IGCAGACIIC

.2340

2280 2220 PCT/US98/04493

8 ႘ ဗ 25 8 S 25 2 5 8 S CAGAAGGCAG TAGGACCTGG TITTITCAGGT ACTGGGAAGCC GGGGGCTCAC TGCTTGCACT CCTCCTCCAC TICCCATGCC TCTATGTTAC CCATCTGTGT CTCCTGTGCA GAAGGAGAGG CACTAGTATA ATTIATAATT ATAACCTATT CIGATTICTT TICAAATATI AGGIGICCTA CACTOTATIAA GCAGTOATOT TOOGAGACCG GGAGGAGAAG GTGGTGGGCT AGTCCTGTGT AAAGCAGGIA TCTTCTGGTT GTCACAGAGT TTCATTGAGT CCAGCTGCAG CCACGTGGCC TOGITAGOAG CIGGAAGACA TICCICCOAC ACITITICCCI TOCIOGCCCA AGAGAGCATO GICTACACGI ICAIGACACA TITICCITICTA AAGGITICAAA GICAAGIGIT TICIGAAGCA TOCCAGACTO AGTOTOCTAT GTCAAAAAAAC TOCATOAAGO TITTIGTGTGA AGATOCTGTT GITIGOCTIATIG AAGGITTIGGO ACTICATICITI GCACTIGITICO CCAAACTITIG GACTIGAGCTA 2) AAAGTGAG TGTTATTCCT AAAAAAAAAA AAAAAAAAACT CGAGGGGGG CCCGGWACCC AWATCGCCSK ATCTOGAGCT GGTGCTATAG GTGACCATCT GGTACATTGA GGGGACCTGT TTGCCTCCTC GIGCTIAGGG TAGGGATGGT AAATATCCIC CCIGCATGGC TITAICCICC CICICATCCC CTGAGGGCAC TCGCTTTGCT CCTGTCAGTA CACACTCCCA AACAGTTAAA CCCAGCTCTA CAGTCTGATT TCTCCAACCG AGTTGAAATT TCCAAAGCAA GTGCTTCTTT AAATGGGGAC AACTOTOCCA ATTTGATCAG CACTCTTATT ACAAACTTGA TAAGCCAGTA TCAGAACCTA TICGCAGAAT ATATTAAAIG TATCCTAAIG GATGAAAGAA CIITTITAAA CAACAACATI NAGGGGCATT AMGAGATGAA GGGTGATTAT GTATTACTTA TCCATTTCTG AATAAACATT CTCCAAGAGC AAGAAGCCAA AGAAAGAAAA ACTAAAGATG ATGAAGGAGC AACTCCCATT ATTCCAACTC TOCAAGAGCT TITAAGCAAA TOCAGGACTT GTCTOCAACA GAGAAACTCA INFORMATION FOR SEQ ID NO: 62: Ĕ (i) SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1851 base pairs 62 2580 2880 2820 2760 2700 2640 2520 2460 2400 240 180 120 420 360 300 600 540 80

5 5 45 55 50 S CATCHATGAA AAAGGAGAGC GAGCTGCCTC GCAGGGTGAA CTCTGCCTCC TCCTCCAACC CIGGOTOTIC TIGCIOGAG GGGGTOTOT TOTGAGCCCT CCCGGTTCTC ACCICGCCTG GODATIGGADA GCCCTIGGTGT CCTGACGGGA GCCAGGTCGG CCTGAGAGCT GTGCCGCTCC GOGIGACTITI CITOTOGCCC CATGGGANGC AGCGIGGGGG CIGICIGAAG GACCCIGCIT CCTCCCCGAG TOGATIOGGAC COGCUTTICT GROTTOTOTT CROCCUTOTS CICTRCTCTC TACGITAACG CCATCGGACT GGAGACCCCT GATTOTOGGA AGGGTTOCCA GGGATAAAGA GCTTCCTCAC CTRIGACCAC GCAGCCCACA GAATICAAAA TCAAGCITTG AGCAGGGGAG TGAGGCAGCC CCCCTGCYGA AGTGGACCCT GACACCATCC TGAAGGCACT CTTCAAGTCC TCAGGGGCCT ACATGAATGG GAATGTTATC ACCTCAGACC AGCCCATCCT GCTGCGGCTG AGTGACAGCC COCCATATAA GAAGIACAAC AACCGGIGIC TOGACGGGCA GCCGATGAAG TOCCAACCTIC CICGACIOGI CCAICCIGGG GIAGCGGAGG IGGIGITITGI GAAAAAGGAC GAIGCCAICA CTCGAGTCAC TGAGGAGGAC ATTGTTGAGC TTTTCTGTGT GTGTGGGGCC CTCAAGCGAG CCAGTOCCTC CAGGCCTCAC TAGTIGGCAAG GGCAGGATGA GGCTGCACCG CTGGGAAGAG TOTOTOTOT CHATGARGET GOOTGGATGG GARACRATTO TORGOOTTOT TOTOCTOTO TITICCIOTAG TAIGITTICTI CAICICAICG CCAAGOTAGG CTIGIGITTI TCAGIGIGIG AGAAGTOGGG GCAGAGGAGG GTGGCTCTGT TTCCCCCAAGG CAÁAGCTTAT GACCAATOGG CCCCATCCAT CCATGACCAG AGGATTATTT TCCTGCCTTG GCAGAGGAGG AGGAGTCAAG GITCACCITY COCTITIOCT IGAGINGIOC IGAANGCOCC ACCCCAGCIC ICITICCCII TOTATOTAAG YTOTTOOCIT GGAGTOOOGT GTOGTOTOOR COCAGAGGAA GTTOTOCAGA CTOGAGOCAT ATATCCAGCT GCCACCAAGG GGCACTGTTT GTTCCCACTT ATGTGAGTGA AGGCTCAGCC TCCCATTGTG CAGTGCTTGG GCACTTAACC ACACCCTOGT TITGTGTAGC CGCCAGCTCT CTTCTGGTTG GGCCTTTGAA TCAAGAGCTC CCAAGATTTG CTTGAGGCTA GCCCAGTGAA RAAAACCAGA GACTCATGTT GGTTCTGGGA GCTCTGCAAA ATCAGTAGCA AGTGCTGGAA AAGGCACATG CCGAAGATAC CCTGCTGGTT GGGAGTGAAG AGAATCCAGG CTGGCAGAGC TGGAGCCAGT TGGGGAGCAC GGAGCAGGGC AGCTCTACCA GGCAAGGTGT TTCCCCAGGA TAGGCGCAGA CAGTTGGGAC COGAGGGGOT GOOTITICCIT TOTGTGTATT AAGCTTTTCA AACAATGGAG CCICAGCCCC AAGCIGATIT CITATCIGGA AATGGTACAC IGAATICICI AGCCCAGGCA GTCCCTGAAT GACCAGGCCA GTGTTGTCAC TGAGTGGTCC GCCTCTCCAT TTCTGGCCTC AGTIGICIAC AGGACAGIOG ICAGGGAIGC GTTTGGAGCT TATTTGAATG GAAGAGGTCA 1260 1200 1140 1080 1020 1440 1380 1320 1800 1680 1620 1560 1920 1860 2040 960 720 660 8 540 480 420

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	ANANGOCOCO GIOTINGCAG ISAIGAGGAG CACACIOING ACACCIOCAI CAGIGACAIG	099
	AAAACAGAAA CCAGGGGGT CCTGACCCCA ACGAGCACTT CTGACAATGA GACCAGAGAC	720
2	TECTCANTIA TIGATICCAGG AACTGAGGAA GATCTTCCTT CECCTGAAAA TAGITETGTT	780
	AAAGAATACC GAATGGAAGT TCCATCTTCG TTTTCAGAAG ACATGTCAAA TATCAGGTCA	840
9	CACCATGCAG AAGAACAGTC CAACAATGGT AGATATGAGG ATTGTAAAGA ATTTAAAGAC	006
2	CTCCACTGTT CCAAGGATTC TACCCTAGCC GAGGAAGAAT CTGAGTTCCC TTCTACTTCT	096
	AICTCTGCAG TICTGTGTGA CTTAGCTGAC TTGAGAAGCT GTGATGGCCA AGCTTTGCCC	1020
15	TECCAGGACE CTGAGGTTGC TITARCTCTC AGTTGTGGCC ATTCCAGAGG ACTCTTTAGT	1080
	CATATECACC AACATGACAT TTTAGATACC CTOTOTAGGA CCATTGAATC TACAATCCAT	1140
5	GTCGTCACAA GGATATCTGG CAAAGGAAAC CAAGCTGCTT CTTGACATTA GGTGTAGCAT	1200
3	GICTACITIT AAGICCCICA CCCCCAACCC CCAIGCIGIT IGINIAAGIT TIGCTIATIT	1260
	GITITIFOTOC TICAGITHOT CCADIOCICT CIGCITGAAT GCCAAGATAG ATTIATAGGC	1320
25	THAATHCTHG GTCAGGCAGA ACTCCAGANG AAAAAAACTT GCANCTICAG TATACTTCCT	1380
	AAAGGGCAAT CAGATAATGG ATATGTTTTA TGTAATTAAG AGTTCACTTT AGTGGCTTTC	1440
ç	AITTAATATG GCTOTCTGGG AAGAACAGGG TTGCCTAGCC CTGTACAATG TAATTTAAAC	1500
ક	TIACAGCAIT TITACIOTOF AIGAIAIGGI GICCICIOTG CCAGITITIGI ACCTIAIAGA	1560
	GECAGATIC CICCGAICGC IGIGGITCTI ATTAICAAAA TIAAGITIAC TIGIATACGG	1620
35	AACAACCACA AGAAATTTGA TTCTGTAAAG AATCCTCTTT AGCTGTGGCC TGGCAGTATA	1680
	TAMATGGTGC TITATITAMC AGAMTACCTG TGGAGGAMAT AMAGCACACT TGATGTAMAA	1740
ć	ATABITICITY TATTITIAIT GACARGACTG ATTGATTGCT ATTCTGTGCA CTTAALTAAA	1800
}	ctgattgtga tgacttwaaa aaaaaaaaa aaaaaaaaa aaaaaaaa a	1851

(2) INFORMATION FOR SEQ ID NO: 63:

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(A) LENGTH: 3542 base pairs (B) TYPE: nucleic acid SEQUENCE CHARACTERISTICS: 3

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STRANDEDNESS: double TOPOLOGY: linear <u>0</u> <u>0</u> (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63

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8 120 .180 GCACAGGCCA CAGAATTCAT TATGAATGAA TATAAGGATC GTAAGATCAC ACGGACCAGC ACAGAAAATG CCATCATGCT GAAACGATTC AATAGGTATC CGCTGATCAT TGACCCCTCT ICCAATGCTG AIGAGCGTCT ICGCTGGCAG GCCAGCTCCT TGCCTGCTGA TGACCTTTGC 8

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PCT/US98/04493

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300 360 420 480 540 909 99 720 780 840 900 960 070 080 140 200 260 320 380 440 200 260 620 089 1740 800 1860 920 980

ITCCIGGATG ACCCTICAG AAAGAACTTA GAGAGIGCAC IGAGAITICGG TAACCCCCTT CTGGTCCAGG ATGTGGAAAG CTACGATCCA GTTTTGAACC CGGTGCTGAA CCGTGAAGTG CGCCGAACAG GGGGAGAGT GCTGATCACT CTCGGGGACC AGGACATAGA CCTGTCGCCA TCCCGGGTTA CITITIGIADA CITICACAGIT ACCCGIAGCA GITTACADAG CCAGIGICIA AATGAAGTAC TTAAAGCAGA AAGACCTGAT GTGGACGAGA AACGATCTGA TCTTCTTAAA CITCAAGGG AAITICAGCI CCGIIIIGCGI CAGCIGGAAA AAICICIACI ACAAGCICIG TOSTITICICA TETTECTIGIE CACEGGGAT CEAACTGICG AGITECCACE AGAICTETGI AACGAGGTGA AAGGGCGCAT TTTGGATGAC GACACGATCA TAACCACTCT GGAGAACCTG AAGAGAGAG CTGCAGAGGT CACCAGGAAA GTTGAGGAGA CGGACATTGT CATGCAGGAG GTGGAGACCG TGTCCCAGCA GTACCTCCCG CTCTCCACCG CCTGCAGCAG CATCTACTTC ACCATGGAGT CCCTCAAGCA GATACACTTC TTGTACCAGT ACTCCCTCCA GTTTTTCCTG GOCATIGCTIC ATCAGGACCA CATTACCTTT GCCATGCTICC TGGCAAGAAT CAAACTGAAG GGCACCGTGG GGGAGCCCAC CTACGATGCA GAATTCCAGC ACTTCTTGAG AGGAAATGAG ATTIGECTIGA GEOCTOGOCTIC CACCOCCAGO ATCCAGGGCC TGACTIGEGGA GCAGGCGGAG CATTOCCTIC AGCOGCATGC CTGCTTCCGA CTCTTCCTCA CCATGGAGAT CAACCCCAAG GACATTTATC ACAACGTCCT ATACGAGAAC CCGAACCTGA AGGGTGTCAC CGACCACACA CAGCOCCTOT CCATTATAAC AAAGACCTC ITCCAGGTGG CGTTTAACCG AGTGGCTCGA COGNIGICA GOLIGABETG CETTCCCGCG TITTAAGGACT TGATTGCAAA GGTTCAGGCA GOGGAGICITI ICAIGICCAT CAIGGAGCAG CCGCTCGACC IGACCCACAT IGTGGSCACA GCAGAAGGCT TTAACCAAGC AGATAAGGCA ATAAACACCG CTGTAAAGTC GGGCAGGTGG GTGATGCTGA AGAATGTGCA TCTGGCCCCA GGGTGGCTGA TGCAGCTGGA GAAGAAGTTG OTOCCTOTOR ATCTOCTCCG TGCGGCCGC ATCTTTGTGT TCGAGCCACC GCCAGGGKTG GACGAGCAAT TIGGCATCTG GCTGGACAGC AGCTCCCCGG AGCAGACTGT GCCCTACCTC CAGGCTTTCC GOCCGATCG CCTGTTGGCC ATGCCCACA TGTTTGTTTC AACAAACCTT GAGGTGAAGC CCAACACTCC TGTCTTAATG TGCTCTGTGC CTGGTTATGA TGCCAGTGGA CATGTCGAGG ACCTTGCAGC CGAGCAGAAC ACGCAGATCA CTTCAATTGC AATCGGCTCT AAGGCCAACA TGCTGAGGAC GTTCAGCAGC ATTCCCGTCT CACGGATATG CAAGTCTCCC AACGAGCGTG CCCGCTTGTA CTTCCTGCTG GCCTGGTTTC ATGCGATCAT CCAAGAACGC TTACGATACG CACCACTGGG GTGGTCAAAG AAGTATGAAT TTGGAGAGTC TGACCTGCGG 2 13 ನ 23 30 35 6 5 S 25 8

GACCCTGAGC CACCTCAAGC GCACCCTGGA GAATATCAAG GATCCTTTGT TCAGGTTCTT ATTEAAATGE CAGATGGEAT GEAGGEGAGA GGAGTTTOTG CAGTGGGTGG AGTTGCTCCC GGGCGCGTGG ACAACGAGTT TGACCAGCGT CTGCTCAACA CCTTCCTGGA GCGCCTGTTC CAGCITICIAC GAGCGGOOTG TOGCAGICTT GIGCACAGAG TAAACTITIC TAGCIGCCCC CAAGCAGACA AACACCGAGA AGAAGGCCAG TGTGGTAACC TTACCTGTCT ACCTGAACTT CTOGTECETO GAGGAGETET GECTGGAAGT CAACGTEACC ACCTEACAGG GEGECACCET ACTOOCAGCT GCATCTOOTO GCGCCAAGGA GCTAAAGAAC ATCCACGTGT GCCTGGGTGG GACCGICATC CAGTGGGTGT CCGACTTCAG CGAGAGGATC AAACAGCTGC AGAACATCTC CCAGCTAGTO AAAGGGATCT TGCCTCGGAG CTGGTCCCAC TACACGGTGC CTGCCGGCAT TOTOGTOCAG GTOTOCOAAG GAAAGAAGAA GCAGACCAAC TACTTOCOCA COCTGATCAA TCAGAGAGAA OTGAAGATOG GCGCAAAGCT OCTTCAGGAC OTTCGCCAGG ACCTTGCAGA CCCTGCCTGG ATGCGGACAC TGCACACCAC CGCGTCCAAC TGGCTGCACC TCATCCCCCA CGACCTGGCC TACGCAGAGA CTGAGAAGAA GACGAGGACA GACTCCACGT CCGACGGGC CGACACCCAG ACGCCTCCT GGCTGGGCCT GCCCAACAAC GCCGAGAGAGA TCCTCCTTAC ACAACCAGGA GTTTCGACAG TGAGTTTAAG CTGGCATGCA AGGTCGACGG ACATAAAGAC TTTCTGTAAT AGTGAAAGTT GGTATTTAAC ATTTATTCAT TTTTAAAATA TTTGGAAGGT CACCOGIGCA GACCICATOT TOACOGIGGA CITOGAAATT GOTACAAAGG AGGATOOTOG CAAGCIGICA CIGICCAAIG CCAICICAAC CGCCCTICCC CIGACGCAGC IGCGCIGGGI OCTOTICOTO COTGAGGOOT ACATCACIGO CACCAGGOAG TATOTOGOCO AGGOCAACAG GTOGACATGA TCAGTAAAAT GCTGAAGATG CAGATGTTGG AGGATGAGGA GGARATANAA CACTAAGCAT GARARAAAAA AARAAACTTA CARNCCNCAA GOTGGANATT GGANGGATAC CAGGAGGTAT TTGGGAAGGC CAATGGCGTG GAAAAGAAAG TOGTTGGTCT ACCITICUCAG TCACGGGITT GAAACTICAA GGGGCCACOT GCAACAACAA GAGGTTGGAG GAAGCTGAAT GGAATCTGAC

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3060 3000

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CTAGTTGTTA CTAAGTANTG CAGTAGCATT NTGGGGAAGA ACA

ATACTAMITY CCIGIGCATC ACACTIAACT CATCTAACTG TICCCCGGAC ANCCICCACT

840 780 8

3

INFORMATION FOR SEQ ID NO: 65:

Ξ SEQUENCE CHARACTERISTICS:

3300 3240

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1541 base pairs STRANDEDNESS: double

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:

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INFORMATION FOR SEQ ID NO: 64:

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2580 2520 2460 2400

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AGCTACAGGC AACACCACTT CCGCGTTTCT CTTGCGCCCT GGTCCAAGAT GGCGGATGAA GOTOTTACAG CAACCCAACG TOTCATOTTO CCATAGTAAA GATGACGGCG COTTGAGGTA CTCGATGCAC TCACAAGCGG GTAACTAGGT GACAAGAAAA CAAAGATCTT ATTCAAAAGA AGGIGATITI AATGATAGGI GICATATATA GGACGGATAA ICIGITTACA TICIGITCIT

240

180 120

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OCCACOCOAC OTOTTOTOTC TOAGATICECO OTOCTOAAGA CTAACOCEGG ACCECGAGAT

360

300

2700 2640

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GACATTCCTA TCACATATCC TACTACTGCC CCAGAAATTG CAGTTCCTGA GCTGGATGGA

540 480 420

TOGTTTGGAA AATGCTGGTA TATCCATGAC CTCCTGAAAT ATGAGTTTGA CATCGAGTTT AACAACAAGA ATOCTGACAA CGATTOOTTC CGACTGGAGT CCAACAAGGA AGGAACTCGG COTGAGITOT GOOTGCAGCG ACTGAAGGAG GAATATCAGT CCCTTATCCG OTATOTOGAG

TOGGOCCAGG AATGTOCCCA AATTTOGACT AGCTCATCTC ATGGCTCTGG GGCTGGGTCC AAGACAGCAA AGATOTACAG GOGTOGCAAA ATATGCCTGA COGATCATTT CAAACCTTTG

660 600

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ATGGSTGGCA GTGGAAATCC CTGATCTGAT TCAGAAGGGC GTCATCCAAC ACAAAGAGAA

TGAAGAATCA AGCCACTGAG GCAGGGCAGA GGGACCTTTG ATAGGCTACG

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TCACCOGATA AGATICCOGTO GTCTCCACTA AAGACCTTAA TOGCCCAGTC CATTTATGGC TCANYTIGCG ATACGGTGGA CACGTGGCTG GATGACACGG CCAAGGGCAG GCAGAACATC

S 8 TGATGGCACC AAGGAGGTGC CCATGAACCC AGTGAAGATA TATCAAGTGT GTGACATCCC GATTICCCTOG AAACATOCCA CCCGOCATAG CCCTCAACAA GAAGAGGAAA ATACCATTTT TARATGGAAG GCCCAGCTGC GCTGTGCTCT CARTAAGAGC AGAGAATTCA ACCTGATGTA THAGGCCTGG GCTGTAGAGA CAGGGAAGTA CCAGGAAGGG GTGGATGACC CTGACCCAGG GOCACGAGGI GOCCICTACC CIGGOCICAI CIGGCTACAC AGGGACTCTA AACGCTICCA

240

180 120 60

90

288

PCT/US98/04493

WO 98/39448

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 883 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 64:

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*	PC 88/39448	PCI/US98/04493	WO 98/39448	L L	PCT/US98/04493
•	586		290		
	The second secon	;	GAAGCTGACC GCACAGTGTC CACACGAATT GGCCCCCAGA AGATGGGGAG TTCTTGTCCTG	Greens	180
	TENSCECCIAS GASTUSMICA TIMALCEANA AICCALMASS INTGETECCT GASATGAGAA	360	COCTITITIONS TOTACOGNAC CTACCACACA GOTOCATICA COCINEATIT	AGATTT	240
2	GGATAATGAT GTGGATGAAG AAGATGAGGA AGATGAACTG GATCAGTCGC AGCACCATGT	420	GOCTICICATT CAMBITICTO GOCCITIGGO COGAANACAG CCAGCITIGG COCTICITIGGO	STIGGE	300
	TOCCATOCAG GACACOTTOC COTTOCTGAA CATCAATGAT TOTOCCATGG GGOCAGOCAG	480	GABACTECTE CAGACCAGA ACCCAGAAG GAGACAGAGC CTGCCACATC CTCCCAGGC	SACGCC	360
5	TOTGGGCAAT TECAOTOTGG GCAACTGCAG CCGGAGGCA OTGTGGCCCA AAACTGAACC		AGGCCCTGGG CCAGGGTGAT TGGACTGAGA ATTTGGCCAC AACCAAATTG ATGCTGGCTG	reacre	420
3	CCTGGGGATG GAAGTACCCC AGGCACCTAT ACAGCCCTTC TATACCTCTC CAGAACTGTG		GAACCAGAGG CCAGAAAGCC TGGCCTTGTC CCCATGTGGG AGCCCTGTCC TCAGCCCTCT	cerer	480
	GATCAGCTCT CTCCCAATGA CTGACCTGGA CATCAAGTTT CAGTACCGTG GGAAGGAGTA	099	TOTCCCCTTG AGCTCAGTGA ATTCCCACCA GGTGCCCACA GCTCCTGGAC TTCAAATTCT	VATTICE	540
15	COOCCAGACC ATGACCOTGA CCAACCCTCA COOCTOCCGA CTCTTCTATG COCACCTGGO	720		TATEC	009
<b>:</b>	TCCCATGCCT GACCAGAGA AGCTCTTTGG TCCGSTCAGA CTGGAGCAGG TCAAATTCCC	780		TAAAAT	099
	AGGICCITGAG CAIAITIACCA AIGAGAAGCA GAAGCITOTIC ACTAGCAAGC IGCIGGAGGI	840			
20	CATGGACAGA GGACTGATCC TGGAGGTCAG CGGTCATGCC ATTTATGCCA TCAGGCTGTC	900 20		*****	07
	CCAGTGCAAG GTGTACTGGT CTGGGCCATG TGCCCCATCA CTTGTTGCTC CCAACCTGAT	096	AND THE PROPERTY AND TH		75
Š	TCAGAGACAA AAGAAGOTCA AGCTATTTTG TCTGGAAACA TTCCTTAGCG ATCTCATTGC	1020			
3	CCACCAGAAA GGACAGATAG AGAAGCACCC ACCGTTTGAG ATCTACTTAT GCTTTTGGGA	1080	(2) INFORMATION FOR SEQ ID NO: 67:		
	AGAATGGCCA GATGGGAAAC CATTGGAAAG GAAACTCATC TTGGTTCAGG TCATTCCAGT	1140	(i) SEQUENCE CHARACTERISTICS:		
30	AGIGGCICGG ATGATCTACG AGATGTTTTC TGGTGATTTC ACAGGATCCT TTGATAGTGG	1200 30	<u> </u>		•
	CAGTOTCCGC CTGCAGATCT CAACCCCAGA CATCAAGGAT AACATCGTTG CTCAGCTGAA	1260			
35	GCAGCTOTAC COCATCCTTC AAACCCAGGA GAGCTGGCAG CCCATGCAGC CCACCCCCAG	1320	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:	,	
3	CATGCAACTG CCCCCTGCCC TGCCTCCCCA GTAATTGTGA ATGCCATCTT CTTCCTTCTC	1380	TTAAGGAATT CGGCACGATC CCGGCAAGTA ACATGACTAA AAAGAAGGGG GAGAATCTGG	reres	09
	TITITITATAA TATIGTACAF AIGGATTITI TIAITGITTA GATTTAACCA GCITTITAAAT	1440	GCGTCGCTCT AGAGATCGAT GGGCTAGAGG AGAAGCTGTC CCAGTGTCGG AGAGACCTGG	CCTGG	120
40	CICTIGITITIC TGTGACAGTG TTAGAAGTTT GTGATTCTCC AAATATGCCT AGATTTAAAG	1500 40	AGGCCGTGAA CTCCAGACTC CACAGCCGGG AGCTGAGCCC AGAGGCCAGG AGGTCCCTGG	CCTGG	180
	ctgatttaat ttatggaaaa aaaaaaaaa aaaaaaaaa	1541	AGAAGGAGAA AAGAGCCTA ATGAACAAAG CCTCCAACTA CGAGAAGGAA CTGAAGTTC	GITTIC	240
45		\$7	TICGGCAAGA GAACCGGAAG AACAIGCTGC ICTICTGTGGC CATCTTTAIC CICCTGACGC	GACGC	00
7			TOSTICTATISC CTACTIGGACC ATOTIGAGCCT GGCACTTCCC CACAACCAGC ACAGGCTTCC	CTTCC	
	(2) INFORMATION FOR SEQ ID NO: 66:	-	ACTTGGCCCC TTGGTCAGGA TCAAGCAGGC ACTTCAAGCC TCAATAGGAC CAAGGTGCTG	TOCTG	420
20			GGGTGTTCCC CTCCCAACCT AGTGTTCAAG CATGGCTTCC TGGCGGCCCA GGCCTTGCCT	TGCCT	480
	(b) Tree: nuclear acta (c) Tree: nuclear acta		cectioaceto etoooooott ecosotetee janaoonen ootoetoote eeteeetting	CTTAG	540
3	TRAUT ::TOTOLO: (0)	**	CCCANGGGAG AGGCAATAAA GAACACAAAG CTGAAAAAAA AAAAAAAAA AACTCGTAGG		_00-
ç	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:	רר	GOGGOCCOST ACCOANTOG CCTIVITOGIG		
	AGAAAATGAA TOTTAGAAGG TGCCTGCGGA GGCGGGACAG AGTGTTTGCT CGCGCTGGAG	09			
9	AAGGCTCTGC TCAGCCGTGA GAGTCCCTTC CTGCCCCACC GATACTGGCA CTTTAAAAAG	120 60			

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1620 1560

1680

1751 1740

CACAGIGATT AACCIGAATG TOCCCITTGA GGICATIAAA CAACGCCITA CIGCIGGCIG AGCGGTGATC ATGGGGGCCC CGGGCTCGGG CAAGGGCACC GTGTCGTCGC GCATCACTAC CTOCTAGECG GCCGGCGCAG GCTGCCGAGC GGGTGAGCGC GCAGGCCAGG CCAAAGCCCT GTTCAGTTAA TAAGTGGTTG ATAAAGTTTC CATATTTTTC TGGAAAAGTT AAAAAAAGTT ACCTITAAAA CATCIGITAG AGCAAAATTA AAAGAGCATT TOGTAGTAAT CTAACTITTT TCTACTGATT TTATTTTGGA TACTAAGGAT GTGCCAAATG ATTCGGATAC TAAGATGCAT GCIGCTITIC CTAAGACTIC TAGTAIGTAI GAATICTIIG AAAATTATAT TACTITTATI TOCTITICCTA CARACTARAG TICCACARAG ARGCCAGRAR GCTICAGITA CICCATGROS AAAAGGGGTG CTGGAAACAT TCTCCGGAAC AGAAACCAAC AAGATTTGGC CCTATGTATA TATCAAQAGA CTAAAQGCTT ATGAAGACCA AACAAAGCCA GTCCTGGAAT ATTACCAGAA CATTGATGAC CTGACTGGGG AGCCTCTCAT TCAGCGTGAG GATGATAAAC CAGAGACGGT GATTCATCCC GCCAGTGGCC GAGTCTATAA CATTGAATTC AACCCTCCCA AAACTGTGGG ACATGROATT TOGAGAAAAT ACGTAATCAG AAATTTTGTGC ATAGATTGAT GCCAAAAAAG COTTIGAAAT CATCIAGIGI GIIGIAIGCA GITAICCICA AAAACAICAG CGAIGICIGA ACACTTOGAG CTGAAGCACC TCTCCAGCGG GGACCTGCTC CGGGACAACA TGCTGCGGGG ACATTICCAG CATIGIGGAA CAIGGIGAGA CACIATATAA AAITICCAGAA AGAAAGCAAC TOGATTTACA GATTTATTOT GAGACACAAA TTCACTOCTG CCTTTACACT AAGAAATOTA ACTIGITICC TITGAAAATA CCIGIGIACI GAGGGITAIG ATTIGIGICA AAAATIGACA ATTITICAATT TOCTATAAAG ATGTATCAAT TAGCATATAG AAAAATATTA CTITAAGAIG INFORMATION FOR SEQ ID NO: 68: GTAACTATTA ATAGTAAGAT GGGCAAACCT CCTAGTCCTT GCATTTAGAA CCAAGGACAC TICCACAGGC AGAAGCCCTA GATAGAGCTT ATCAGATCGA COGCIGOCCC TICATGAGCT GAAAAATCIC ACCCAGIATA GCIGGCIGIT GOCGTOTTAG CCAAGGCTTT CATTGACCAA GGGAAACTCA TCCCAGATGA coerecease ereveneuse oscenioses acarecases oseriacions SEQUENCE CHARACTERISTICS: ATATATOCTO TATTIATTIT GICGITAAGC ATACITICAG TITACICAGA SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 1751 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 1020 1500 1080 480 420 360 300 180 120 720 660 600 960 900 780 23 20 330 15 5 \$ 6 8 S 50 TRAGTOCTIT TACHAGCACC ARAGITICAAT GARTITITCAA CARRATGIRA ITARAGICIA TITIGATICCA AATGIGIGAT CIGCCCIGAT AAATAACAAG TIATNOTACC AICTCCCCCG TOTTTTCAGT TATGACTCAG GTTAAGAAAT GTGTTTTAGG ATCTACTTGC CAGCTRARAA TIGARARCAA AGRICIOGAC RACARRACAG CCARAGGTOG GGGTCRAGAR GCCACGAGAT TATGTATTAA AATGTTTTTG AATTGTGAAA TATTAGAATA TIGTTACTAT CCAGGAAAAA CTGCAGAGAG CCCCAGTCTT CACCTCTGGT TGACCATGAG CTCTGTGTAA OCTOTOLOGI: GTACCINGOT GTAGAATGOT ATGCACACGT GCCAGGTGTA GTGTGCATAT ATRATCAGAT GCAGTATCAC AGCTGTGTCA GACTCTAGTA CCAGTTGGGC AATCAAGGCA TIGACCCAAC ICAAAAICIC CAIGGGAAAA TACCIGICGA TACCCACAGI ATIGIIGAAA 5 NAATAGGNAG T CCAATAAAAA AAAAAAAAAA AAAAAAAAC TCGAGGGGGG 3 CCTACTCAGC TGGACTTAAA AATAAAAA ATCOTOCAAA GCATTAAGGA AATCITGTTA CIGCTAAGIG TIGCIGACCC AGGAACAACT GCAGGAAGTG GATTOCAGAT TIGCTIGIGI CCICAGGIGA IGATGAGGGC IGITITICCCC IGITIGICCIT TACATAGAGC AAAGAGAAAT TICCAGAATI ICTARAATIC TGGAAAGAGA ATITICCIGA TECTEACACT CATECTTECT CIECTAGAGT GICTEGTING CATGATCATE TECTACETAG INFORMATION FOR SEQ ID NO: 69: INFORMATION FOR SEQ ID NO: 70: Ξ Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70: SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: AAGGCTAAGG CAGATTTAAG CTCTGAAAGC ATTCCACAAC ATACACACA SEQUENCE DESCRIPTION: SEQ ID NO: 69: (A) LENGTH: 245 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear **909** (A) LENGTH: 508 base pairs TOPOLOGY: linear TYPE: nucleic acid STRANDEDNESS: double OCCCGGTACC CAATTCTCCG

> 240 180 120

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480 420 360 300

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GGATGGTTTT

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180 120

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TATGTTAACC

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AGAAATGTGT

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GGTACCCGCG

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5		PCT/11/598/04493	MO.	WO 98/39448	PCT/US98/04493	83
•	293			294		
	CCATTICTIT CACTGAIACA AGGAAAACTG CAGGGTTAAA AAAAAAAAA AAAAAAAAA	240	J	GICTITICCCT GITAGGAITT TAATAGCAGA ACTGTATGAC AAGITTAGGT GATCCTAGGA	540	
	NGNCO	245		tatgitaaat tcaaattaat giaaaacaga ttaacaacaa caaagaaact gectatitga	009	
S			'n	GIGAAGTCAT GCTTTCTATT ATAATAACTT GGCTTCGGTT ATCCATCAAA TGCACACTTA	099	
	(2) TAMPORMARTEN FOR SEC TO NO. 71.		-	inctottatc tgattgttta taataaagaa tactgtactt ataaaaaaa aaa	113	٠
9	(i) entitating cutabamparence.		01			
2	(A) LENGTH: 151 base pairs					
	(B) TYPE: nucleic acta (C) STRANDEINESS: double					
7	(D) TOPOLOGY: Linear		5	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 862 base bairs		
3	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 71:		:			
	ATGITECTEA TEAGGATECA CITISTECTIC TECAAGTATT CCTCCACCTT CATAGTEACT	09		(D) TOPOLOGY: Linear		
20	CCCACCAGCA CCAGCAATAC AGCTAGCTAC CTGTGGCCTT GGATCTCAGC CACCATGGCT	120	70	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:		
	GOGAGAGGGA GCAGCTGGGC ATGTACCCTA AATGCTGTTA CCAGGGAAGG ACTCCCAGAG	180	Ü	GAAAGTCAGA GCTGTCCAAT CCCTCAGCAC CTTTTAGATT TGCTCCAAAT TAGAAACGTG	09	
;	TGAAGACAAG TAGGGACTTC CTGCAGAGGT GGTACATGTG CTCTCTGTAT CCATACTTTT			GOGACTATOT GITCTGGGCA ATCACAGGTC TGGAAAATGG CTCTGCAGGC TCTTGATAGT	120	
23	THITHTH THIGAGALA GAGTHICACC CHISTIGCCC TGCTGGAGT GCAATGGTGC	300	3	GAGACAGTOG TCATCTTACC AGACATGCAT CTGATTTTAA GCCTCAGGCT AATCCACAAT	180	
	GAITCHCAGCT CACTGCAACC TOTOTGCCTC COGGSTTCAA GTGATTCTCC TGCCTCAGCC	360	9	GCTCGGCCAT GCCTATGATT AACAAACAAA AGCAAAATCT GCTTTTATAG TTTAGGAAAC	_340	
30	· ·	361	30	ctogatagaa cagtatttit cagcattett ggataaagca sttetgeatt tttaaattgs		
			O	gactecagaa Gtgactetet atagttgtga aatacaaaaa atggtatgtt tgatcagaaa		
;			* v	AGGAAGCCCG TGCCTGGCAC TTGGAAAGAT ACTGAGCATC ATAACCCTAA TGAGAAAATG	420	
33	(2) INFORMATION FOR SEQ ID NO: 72:			TAGGCTCTOT GARIGITAAC TACAAATCAG GITAGGAAAG CATAIGACAC CCTITOTCAA	48	
	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 713 base pairs		~	ACTANGCTTC ACTROGRAGA CCTOTOCTCA TRGAAGAATA TOCTTTIAAAA GTATCAATTT	- 240	
9			40	TCCACAGTCG ATGATGGAGA AAAGTTCATT TGCACCAGAA TGCTGATAGT CACAATACAC	009_	
	(D) TOPOLOGY: linear		•	ACCTIGACAT ATATAACAAT ACAGTITICT GTAAACAGAA GTICTICCTC TICCAAITICA	099_	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:			CGAGTCAGTC AGACCATAAA TATTGCATGT TTCACTTTAG AAACTGATTC ATTTTAGAAA	720	
45	AGGATCACAC AATAGAGAAC ACTGTAGTAA CATTTCGGTC TGCTCACAAG ACCCAGAACA	09	<del>ડ</del> ૦	GCAGATCTIGG ATTATTTTTIGG AGGGTAGAAA TGAAGGCTAT TTCTIGGCATT CTTGCTCAAA	780	
	TIGAICAGIT TITICITICITIC CITTATTATT TITICICITAA AAAATTICIGA AAAGITICIT	120	~	AAGTCAATAT AIGTACATTA AGTATAAAAA AGGGTCTCTT TCACCTCTTT TGTTTCGTAG	840	
5	THAGCHAGAT GATATTTTAA TAGCHGGGAG TGCTTTGGAA CTAINAAGAT GTCACTACTT	180	20	CATTGGCTAC ATAACTCGTG CC	862	
3	AACACACATA CCTTATIGTTT TGTTTTTACAC TCAGTATAAA TCAGGAGAAG	240				
	TTAGGCAACC, ATCTAGGATT TAGAATCCTC TTTTTATTG TCTTCTAAGG ATATGGATGT	300				
55	TCCCATAACA GCAACAAAAC AGCAACAAAA ACATTTCATA AATATCACTT GATAGACTGT	360	22	(2) INFORMATION FOR SEQ ID NO: 74:		
	AAGCACCTGC TTAACTTTGT GTCCCAAATA TTTAGTGTGT ATATATAT ATATATAC	420		(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 4602 base pairs		
9	ACACACACAC ACATATÁTAT TCAACAAATA AAGCAAAATA TAACATGCAT TTCACATTTT	480	09			
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CCCAGGGGGG GKGGGGAGCA GCGCCGARGC CGCCGCCTCC GCCTCCGCCG CCTAGGACTA GATTETECCC TECCTEAGGA AGITTECTATT GETGCATETA GACCTAGECG GGGCTGGCGT OCCCCTUAG OCCOTOCOG COGCOCCTOC AGGAACCACT ACCAGCCGCG TOCTGAGGGG GCAGCAGGAC CAGGAAGGGG GAGAGGCGGC CAAGGCGGCT CCGGAGGACC CGCAACAACG GATOTTOGAG AAGAGCATCA GTCTCCAGGT GGCATTAGTA GTGAAGAGGA AAGACCOGTT CATTGCAGCT CATTTGCAAG TCAGAACCAA ATACAGACCA ACTTGATTAT AGTAGTAGGA CATCTGTTTC TCGCCATCGT GATACAGAGA ACACCCGAAG CTCTCGGTCC CTATGCATAG AAAAAGTAAC AACTGATAAA GATCCCAAGG AAGAAAAAGA GGAAGAAGAC GEAGEAGECE AGGEOGEGAA GTECCEGGTET CEAGTTEAGG GEAAGAAGAG TECGEGACTE AGGCCGAGTA TCCCCGGCGG CGAGGAGCAG CCCCAGGGCC AGGCCTCCCG ACGTCCCGG датеррал саловества ссостветке соссостать сластвтоге ссоссозлал GOGGOTOGGG GACGGACAAG CCCCGATGCC GGGGGAKACG GAAGAGCCGA GACCCCCGG TACCAGCATT TOCTGAAGAA GAAATATGTA TGTCCCCATC CCTCCTGTGG ACGACTCTTC GAGAATGAAA TIAGAGAGGA TGAGGAACCT CCAAGGAAGA GAGGAAGAAG ACGAAAAGAT ACCTACAAAC CCCACTTAGA AAGGGAAAACC CCAAAGCCAC GGAGAAAATC AGGGAAGGTA GAAGAAGAG TOTTAATCAG TGAAGAGGAG ATACCATTCA CACACTOSCO AGAAGCATTA CAATOTGAGA TCTOTGGATT TACTTGTCGA CAAAAGGCAT TOTGAATATT GIGCICGGGC CITCAAGAGI ICCCACAATC IGGCAGIGCA CCGGAIGATI aaagaagaga aggagaagaa ggaaattaaa gtogaagtag aggtogaggt gaaagaagag CTCTTAATTO GCACATGAAG AAACATGATG CAGACTCCTT CTACCAGTTT TCTTGCAATA AGGCTTCAGA AGCAACTTCT GCGACATGCC AAACATCATA CAGATCAAAG GGATTATATC ATATETTOGG CACTAACCCA GAGTCCCTGA CGCAGCCTTC AGATGGTCAG GGTCTTCCTC CTGAGGTGCT GATTGCAGAA GCTCTGGCTG CCAATGCAGG CGCCCTCATC ACCAGCACAG TCTOTOGCAA AAAATTTGAG AAGAAGGACA GCOTAGTGGC ACACAAGGCA AAAAGCCACC GGATGICAAA GICATACIGC AGIGGGACGG AACGGGIGAG CCIGAIGGCI GAIGGGAAG TICTICCIGA GCCCTIGGGA AACICAACCI CIGGAGAGIG CCIACIGITA GAAGCIGAAC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74: GATGTGGAAC TGTCCTTGCC CATCCTCGCT ATTTGCAGCA CCACATTAAA CACCITIACC CAAAAGGAGA AAAAAGCCIC CAATCCAGIA IGICCGIIGI AAGATGATCC AAGAGATGAG AGAOGAGGAG 1320 1260 1200 1140 1080 1020 1500 1440 1380 840 780 720 660 600 540 180 120 1680 1620 1560 960 900 480 420 300

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OTCAGAAATG CATGATGGCT CTTGGAAAGA ATGACGTTTT GCTGGAAAAA AAAAAAARAA

ATTECTATES TOTOTOCOTO AGAAGOTAGA ATCGAAGGOT TACCOTATIO ATTGTTATI ACAAGGTCAT ACCOCCAGAA GCCCCAAATC CTATTTTOGC TCATCTTCAG GTAAAGAGTA TIGIGITIGGG GAATAGATIG CITATIOCIG TICACIOGAG AGAAAAGGIA GIGITITITOT

CMSTTTGTGT TICACAAACA TOGCTTATCA ATTTTTTCAA AGAATTCTTT TTTCCCCAAAA

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AATAAAGATA GAAGGGATGT AGAATATGCT TTCCTGCCAA CATGGTTTGG AGTCGACTTT

TGTTTCTTTT CAGAGIGITG CAGATITGCC ATTICICCAT AATATOOOGA TAGAAAATOO

CICCICICAG GICCCTITIA CACTITITGA CIAACIAGCA ICIATATICC ACACTIAGCI GGTATATTGA CTAGATTTGA AAATACAAGA TIGATTAGAT GAATCTACAA AAAAGTTGTC

> 3480 3420 3360 3300

TITGGGATGC CAGGGAACAG AGAGTGAGAC ACCTACAATC ACCAGTCTCA AATGCGCTAT AGAGGAGTAA CAAAATGTCA TTTCTGAAAG AGGCTTACTT TATACCAACT AGTGTCAGCA

> 3240 3180 3120 3060 3000 2940

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TAATAATTOC TITIGGCTTTC ACCIAAAATT CIGGGCATCA CAATTICCTT GGGATAGAGG CCCCTTTAGT AACACTICTG AAGAGGAAAA ACTICAATAG CCAAAGITAA TAATCCTATA TAGTGAAGGA ACAAAGTCTA TGAGTCCTAA AATTTTAAGT CAAAGAAAAC TGCTCTGTTT TOTTTCTTOC TTTOCTTTAT CAGTTCATTC CAGCICCCTG TTAGTGAAGG ACACTGCTGT TITTIGICCC TACCATTICC TIACATTICC CTIGGGGCCC AICTCIGGCI CCTIGCTITT

2880 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2220

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GTTCAGAATC TAAATTACAG ATAGATGATT GTTTCTTGTG AATTTGTTTC TTTTCCTTTT AUGCTOTTIG AGATTITIAC AGIGTIGAAA CITAAGAAIT TIGAGAGGGI GAGGAGGGIT TAGCCTAGIG CTITTTIGGA AGCCTITTIA GGGAAGAAIG TIAGGIICAI GGTAACIAGI CATAGOTOTO CAGTOGOTTT TAAAGAAAGC TOGTOCTCAG CACTAACAAA ATCACTACAA GOCTOCTACO COCTGOGGA CACTTATOTT CAAATACCAT AGAATTOTAA TOTOTGAAAT

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TAATCATATA TIACIGICIT CIAAAICCCI ICICCICCIC TACIGCIGCC CIAIGGIICI TOGGCAGATO COCCTITICCA ACCIGIAACT CIGATGIGCI CIGGATCAGC TITTAACTIT GUICTIAAAG TOAGGOTTAT TUTCATACTU GOTTOCAGCU ATCAGCAGAC TICCTGUICA TOCCCTTATA AAATTGATGT TGTCTTTACC AGAAAGGTAG ACAAAAAAGA AGCAGCAGCA ATCACAGCAC ACACATACAT ACACCCTCCA CCTCCCCATC CCCTGTTCTC CCTCTGTTGC

2160

2040 1980 1920 1860 1800 1740

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AAAATCTAAA GCATTTAAAAA TCTAGTGAAA TAACTGAAGG GCCTGCTCTT TCCATTGTGG GAAGACTTOG GGCATGGGAC AGCTCAGACT TTGTATTTAA AAGTTAAAAA GGACAAAAAA TETTTOTOGO AAGCGGCAGC AGTGGAGGCA CTGAAGGGCT GGTTATGAAC TCAGATATAC

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TOGOTOCTAC CACAGAGGTT CTGATTGAAG

ATTCAGACTC TGCCGGACCT TAGTGGACAG

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(D) TOPOLOGY: linear

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	TITITICICAC ACTIVICCIT TOTCICCOTA ANTITICALITI GCAGIGOTTA GICATCAGAI	3540	GCCAAAAGAT ATTTGACGGT TTCCAAAATT CAGATTCTGC CTCTGCGGGAT AAATATTTGC
	ATTTTAGCCA CCTACACAAA AGCAAACTGC ATTTTTAAAA ATCTTTCTGA GATGGGAAAA	3600	CACGAATGAG TAACTCCTCT CACCACTCT AAGGTCCAGA CAGAAGGTTT TGACACATTC
2	AATGTATTET CETTTECTAT ACCOCTETEC CAACAAAAA ACAACTAGTT AGTTETAETA	3660	5 TIRECROTEA ACTICATIONS GANCTIAGRAN GANCTIOC COCYCTIGRAN GARDANCCIO
	ATTAGAAACT TGCTGTACTT TTTCTTTTCT TTTAGGGGTC AAGGACCCTC TTTATAGCTA	3720	TERMERICA GECTIONISC AGGINATIT GARGESTACA ATCAGANGCA AAAGCTICTTG
9	CCATTIGCCT ACAMAANT ATTGCAGCAG TITIGCAATAC TAAAATATTT ITTAIAGACT	3780	10 constraint thanastar constraint extensions accounted accordant
2	THAINITIT COTTITION AGGGARGOT GCARAGRAGA GRIGOTIGRA TRAACTATO	3840	
	TCAGCCGTIT CCCTGCTITC CCTTCTGCTC CATATGCCTTC ATTGTCCTTC CAGGGAGCTC	3900	TOTAL CANCELLA CANCEL
15	TITITARICIT ARAGINCIAC ATTICARGCT CITAGOCAAA TICIGITACC TITITAARAA	3960 1	15 TATAGACTET ATCHARACC ACAGETICATC CCCAACCCTA GECTECGAA TGETAAGATET
	CTCTTCCCAC TOCATATTIC CATCTTGAAT TOCTGCTTCT AAATTCTGAA ACTGTAGTTG	4020	TEATHABAITET AGABARTETA THERIPACAAT TACAGARTIC ABATTATTIGGA ABAGGAITOTIG
ć	AGAINCAGCT AITHAIRTI TCTGGGAGAI GIGCAICCCT CTTCTTIGIG GITGCCCUAG	4080	20 negeritere eccessence ecteritees Aceaecades reconterer
8	CITCITITIC CIAACIGAGA CICCITIGAIA TOCITICAGAG AAITIAGGCA AACACIGGCC	4140	
	ATGGCCOTTOG GAGTACTTOGG AGTAAAATAA AAATATCGAG GTATAGACTA GCATCCACAT	4200	ACARGEMEN COSTISTIBLE CANCENTER THEFTICALTY AND APACTACAG CITICS ACAR
25	AGAGCACTTG AACCTCCTTT GTACCTGTTT GGGGAAAAG TATAATGAGT GTACTACCAA	4260	25 mmmmme 2.5 mmmme and memory of the manufactures of the manufact
	TCTAACTAAG ATTATTATAG TCTGGTTGTT TGAARTACCA TTTTTTTCTC CTTTTGTGTT	4320	ANAMANGO MACAMANA MACAMANA MACAMANA MANAMANA MANAMANA MANAMANA MANAMANA
	TITCCCACIT ICCANIGIAC ICAAGAAAT IGAACAANIG TAATGGAICA AITTAAAAIA	4380	ALTERNATION INCOLUENT INTERIOR INTERIOR AND ARTHUR AND
30	TITIATITCT TAMAMGCCTT TITIGCCTGT TGTAATGTGC AGGACCCTTC TCCTTTCATG	4440	
	CCAGAGACAG GTAGTTACCT GAATATAGGT TGAAAAGGTT ATGTAAAAAG AAATTATAAT	4500	AUTHORISM AUTHORISM AUTHORISM NOOCOCOCO CECEGOORISM NEWSTANDS
35	ANANGGANA CTTIGCTTTT CAAATCTTTG TTTTCTCTTA TTCTAGGTAA GGCATHTTAA	4560	35
	aaataaatat gtaaagaaga aaaataaaag ttgicttcat gg	4602	(2) INPORMATION FOR SEQ ID NO: 76:
Ę		4	(i) SEQUENCE CHARACTERISTICS: 40 (A) LENOTH: 475 base pairs
}	(2) INFORMATION FOR SEQ ID NO: 75:		(B) TYPE: nucleic acid (C) STRANDENESS: double
Ý	(i) SEQUENCE CHARACTERISTICS:	4	45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:
4	(A) LENGIN: 1233 DOME POLLS (B) TYPB: nucleic acid		
	(C) STRANDEDNESS: double		GOCACGAGAG AAATGTTIGA TICTCTTICC TATTITAAGG GAICTICICT CITGTIGATG
	(D) TOPOLOGY: ATREAE		TIGAAAACTT ACCTTAGTGA AGATGTGTTT CAACATGCTG TTGTCCTTTA CCTGCATAAT
20	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 75:		3U CACAGCTATO CATCTATTCA AAGTGATGAT CTGTGGGATA GTTTTAATGA GGTCACAAAC
	CGCGCCCCGG GCCGGCGGGT TTCTCTAACA AATAAACAGA ACCCGCACTG CCCAGGCGAG	09	CANACACTAG ATGTANAGAG NATGATGANA ACCTGGACCC TGCAGANAGG ATTTCCTTTA
ž	CETTGCCACT TTCAAAGIGG TCCCCTGGGG GAGGTCAGCC TCATCCTGAT GATGCTGCCA	120 5	55 GIGACTOTIC AAAGAAAGS AAAGGAACTT TITATACAAC AAGAGAGATT CITITIAAAT
રે	AGGCGCACTT TITATITITA TITIALITY ALTITITITY PAGCATCCTT INGGGCTTC	180	
	ACPCTICAGAG CCAGITITITA AGGGACACCA GAGCCGCAGC CTGCTCTGAT TCTATGGCTT	240	CICCITINICA CIVICAGGIG GAAAGATITIT GAAGIGITITI AGAATCADA GITCITIGIGA
9	GOTTOTTACT ATAAGAGTAA TIGCCTAACT IGAITITITICA ICICTITIAAC CAACTIGIG	300	09

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AACCTAACAA GATTATCCCT TCCTAAGAAT ACTTAACCTT CCTACCAAAT TAAAA

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	50		45		<u>.</u>	à		35		į			25		2	3		15		-	5		5		
(2) INFORMATION FOR SEQ ID NO: 79:		CHACCTOTT GOGAATTACA CATRICAATA TAAACAAAAT ATAAAGT	5 ANGANGTAGC ATATOTGAAC TATAATGTAA CAGTGANTAA TTTGTNAAGT TCGTATTTCC 1860	ATTGAGCCAC AGITGAGCTT GGAAAGTTTT TGTCAAATGC NGCAAGAGAT AACTCTTTTT 1800	тттопанта осолавська тестттова тараантваа твасальное аталаасттт 1740	CAACTETECA CCATCAATGT AACTECATGG ACATTGCTGC TETTGGTGGT GTTATETAAT 1680	ATTITITIET TITITICITY TOCCICCITI ANDACCITIG GRACATIGG AATACCCAGC 1620	AGAAAATGTC ATGTGATGTC TCTCCCCAAA GTCATCATGG GTTTTGGATT TGTTTTGAAT 1560	ANGTONAATT GTACTTGATC CTGCTGAANT ACATCTGCAG CTGACAATGA GAGAAGAAAC · 1500	GATTGCTGCT AACAGTCAGG GTACAAITAA GGTGCTAGAA TTGGTATGAA GGGTTAACTC 1440	AMATGAATTT GITHGTOCTG TGTGCTGGAG GGCACTACCA GATGGGGAGT CCAATGTGCT 1380	AACTITIAAG TITGATACAG TCAAAAGTOT TCTOGACAAA GACCGAAAAG AAGATGATAC 1320	TTGTGGAAGT GAAATAACT CTCTCTACCT GTACTATAAA GGACTTTCTA AGACTTTGCT 1260	CANGGGTCAT ATCAATGAAA AAAACTTTGT AGGCCTGGCT TCCAATGGAG ATTATATAGC 1200	CTCAACAGAC AGTCAGCTBA AACTGTGGAA TGTRGGGAAA CCATACTGCC TACGTTCCTT 1140	асандаесот анасенотет сттипосная стттопаног сотпасания ттотететос 1080	AGATCACTOT GTCCACTACT ATGATCTTCG TAACACTAAA CAGCCAATCA TGGTATTCAA 1020	TANTOTOTOC TOTOTTANAT TCHOCCCCTC TTCCAGATAC CATTTOCCTT TCCOCCTOTCC 960	AAAAGTGAAG CTGTGGTCTA CCAATCTAGA CAACTCAGTG GCAAGCATTG AGGCAAAGGC 900	TTGGAGIGIT GACTITAATT TGATGGATCC TAAACTCTTG GCTTCAGGTT CTGATGATGC 840	ттрапоская сантелем саемалесте аласстетат смежеское асамаское 780	CAGTIGGAGT AGTIACCATA AGAACCTGTT AGCTAGCAGT GATIATGAAG GCACTGTTAT 720	TOCHOTOGAT ATTUNTINGS CTOHGRATCH ANTGROCTOS ARTICGRANA TCHOCTOTAT 660	посомпост осмотрасма моммантим мотетитома титеменето темпесносы 600	TCTCTMTAAT GGTTCCAGTA TAGTCTCTAG TATTGAATTT GACCGGGATT GTGACTATTT 540

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S

GGAAGAAATG AGTGGCTTAT ACTCTCCTGT CAGTGAGGAT AGCACAGTGC CTCAATTTGA

AGAGCAACTG GAACAGATCC AGAAGGAGCT AAGTSTTTTG GAAGAGGATA TTAAGAGAGT

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S

GTCTCGTATC TCAGATGACA GTCGAACTGC AAGCCAGTTG GATGAATTTC AGGAATGCTT

AGETICETTET CENTENCACA GINGINITAT TGATTCCACA GANTACAGCE AACCICCAGG
TTTCAGTGGC AGITCTCAGA CAAAGAAACA GCCTTGGTAT AATAGCACGT TAGCATCAAG
ACGAAAAGGA CTTACTGCTC ATTTTGAAGA CTTGGAGGAG TGTTACTTTT CTACAAGGAT

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(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1168 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

GTCCAAGTTT ACTCGATATA ATTCAGTACG ACCTTTAGCC ACATTGTCAT ATGCTAGTGA

120 1120 1180 240 240 300 360 360

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:

5

ACATIGCAGEC CAACTACAGA TYCTTATIGGA ATYCCTICAAG GTTGCAAGAA GAAATAAGAG

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:

30

GCTGCACCTT TCAAGGACAA AGTGCTGGTG GCGGCGCNGA ANGAA

120 180 240 300 360 420

TRATTECTOS COSTOSTANA CARCONSTAC GTGANCOCOC AGACCANCOT GACOSTOTOS AGACTICTOGA AGAGTAGOCO CTITOGACGANC AGCTOGACCA AGAAGGACGC GCANGACCTIC GTGAGCGTOC CGACGGAGAAM CTCGARGACT KOGCGCCGA CACGCGCTTC TROGTGAGCC AGCCCGGAGAGAC GCGCCCTGGG TCGCCCTGGT GGTCGTGAGG

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(2) INFORMATION FOR SEQ ID NO: 78:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1907 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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GENERAGEST TOUTCOCCCT GACCETOTICS GTGCCCGGGG CCCGGGGCCG GGCTCTCGAG

TICTCTOTOC TCTTCGACTG CACCGCACTC GCGCGTGACC CTGACTCCCC CTAGTCAGCT CACCGCGCGCG CGAACCRGCG TCGGGGCTCG CGGCGTGTTG

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:

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(2) INFORMATION FOR SEQ ID NO: 77:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 465 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

5	WO 98/39448 . PC	PCT/US98/04493	WO 98/39448
	301		305
			CTGCAGGACA GTTTGGTCAG AGCTGCAATT CTTAGTCCAT GGTCTAATGC TTGAGTATCT
	GCTGGGGTGT CCCCKCSGCC ACCATCGTCA TCGCTTACTT GATGAAGCAC ACTCGGATGA	09	CTICITICCC TITICCIONCE CAGAATCAG CIGAGAATTC ATTCGATIOF CATGCCICTA
v	CCCATGACTG ATGCTTATAA ATTTGTCAAA GGCAAACGAC CAATTATCTC CCCAAACCTT	120	ACCOCUMACT GRANTINGTH GOTHOCACTH TAGTICTAGA ATCACCTGTH
`	AACTTCATGG GGCAGTTGCT AGAGTTCGAG GAAGACCTAA ACAAGGGTGT GACACCGAGA	180	GACTOCTORS ACTITIONAL CICILITIES GAGGOTICIC ATTICCCOCT
	ATCCTTACAC CAAAGCTGAT GGGCGTGGAG ACGGTTGTGT CACAATGGTC TGGATGGAAA	240	A ANTHONORY OF THE ANTHON CONCORDED ANTHONY CONTRACTOR
01	GGATTGCTGC TCTCCATTAG GAGACAATGA GGAAGGAGA TGGATTCTGG TTTTTTTTCT .	300 10	ANTICULOR CONTINUES CARRESTOR DESCRIPTION CARRESTOR CARACTERICS ANTICARTIES
	TICITITITY TITTIOTAGYY GOGACTAACY TYCTGAATGG AAACAAACYY GYYYAAACAC	360	TOTAL TOTAL
;	TITATITITIA ACAMOTOTAA GAAGACTATA ACTITITGATO CCATTGAGAT TCACCTCCCA	420	ATCHARAMIC CHEATLICE NONDOMERIC CICICIANO TOTORNOCO TOTORNOCO CONTROLOCO CONTROLOCO TOTORNAMAS
2	CAAACTGACA AATTAAGGAG GTDAAAGAAG TAATTTTTTT AAGCCAACAA TAAAAATATA	480	CHRAINSOCHE THRESTABLET ANAUTOTICS ANGTHURAT GARCICOCCAT AND SECTIONS
	ATACAACTIG TITICICCCC ITITICCITIT AAGCIATITIG TAGAGITIAT GACTAAATAG	540	GCACCITIVES CINCIPIES AND STRAIGHTAND GAGAAAGAC CEATAITRIG AGTCAAUCTA
20	TCTGTGCAGG TTCATAGACC GAAGATACTA CACACTTTAA ACCAATAAA AAGAACCAAA	600 20	CHACCHARTER MICELANDICA ACCERANITIC CACATABEDA CACTABEDA CIVITICIONE
	ACTIVATIVGA AAAGACATTG AATCACCAAG GCCTGGGATC AACCTGGGGT GTCCACAGA	099	ACTACTOCA CANGADAD CADACTOC CATAVADA ANTIACTICS TGACADOTTIC
ž	AAAACAAAAA CCCAACCAAA CCAAGCCCTG TTGTGCTCAC TGGTGCAAAG AGAAGATCAG	720	CHARLAGE ATTRACTERS ADMITISAGE CACCATCATT AAACAGTTAT GTIGITCIC
3	GGCAGCITAA GTGGTCTAAG RAICCTTCAG GCATTCTITA AGGAGAAAA GGATACCTTT	780	THE PROPERTY THREE PROPERTY CHIPCOCTING ACTIVATIONS ACTIVITIES THREE TRANSPARTS
	GATITIGICA GITICARGCT CROGATITIT ITITITITIC CITCICAGGG TITAAGAGAI	840	AACTICOTOR DEADURE TITICAAATIG CICILCOTICOS COAAAATAIT GAAATGAGG
30	TITITITICAA ATAGTGAGGA ACTGACCATT ATATGCCTTC ACTGGCTTCT TGTGCAATAA	900 30	AACTICATICA TAAACTICTIA CAARTAAGCA GTGTICCCTAA CAAGCAACAC AGTAATICTIG
	TATGATGTTT TAAGTGTGCA AACAAGTTAG AGCTGGCAGC TGAATGATAG ACAAATAGTG	096	ACAMEMINA OFFICE AREA OFFICE OF SAME OF TATABATICS OFFICE OFFICE
;	CARATTIOCC AGCTIGGAGA TAGAAAGGAA TICAACAATA TAICAAATAC TITCCTICCC	1020	NATIONAL STATEMENT OF STATEMENT OF STATEMENT STATEMENT PROPERTIES OF STATEMENT STATEME
32	ACCITITINC TITITITITI ITITITICIDA TITICALTICIS GITACAGIGO CALABACCIT	1080	WEINTILL CONTINUE WORKSTOND INTEGRALIA STREET
	GTTACATATG TATATCAGAA TGTAAGAAAA AAAATTTAT TTAAAATAT TTTTGGCAAA	1140	ANTALGANAA TAKTANAKAT TUMAG
40	AAAAAAANNA AAAAACTUGA GGGGGGCC	1168 40	
			(2) INFORMATION FOR SEQ. ID NO: 81:
45	(2) INFORMATION FOR SEQ ID NO: 80:	45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1290 base pairs (B) TYPE: nucleic acid
	(i) SEQUENCE CHARACTERISTICS: (A) LEXGTH: 1285 base pairs		
50	(B) TYPE: nucleic acid (C) STRANDENESS: double	20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 01:
	(D) TOPOLOGY: Linear		TCTCCAGCCC CAATTICTAC GCGCACCGGA AGACGGAGGT CCTCTTTCCT TCCCTAAGGC
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:		ACCATGGCT COTGGTCCCA AGABCATCT GAAGCGGGTG GCAGCTCCAA AGCATTGGAT
25	AGAAAATCAC ATOCTAACAA AGAAGICTIGT CTAAGACAGT ACATCTOCTG TTGAACTTGC	60 09	GCTGGATAAA TTGACCGGTG TGTTTGCTCC TCGTCCATCC ACGGTCCCC ACAAGTTGAG
	ATCTITCCAC AGGACTITCT GITTITIAGGG ATGAGACTAT TCTCTGCTTC ATCAAGGAAA	120	AGAGICICIC CCCCICATCA TITICCIGAG GAACAGACIT AAGIAITGCCC TGACAGGAGA
9	GAGANATOTT CAGGETTGTA GGGATGGCAC ACTIVITAGT TCTGCCTGTC TGANAGGTTC	180. 60	TGAAGTAAAG AAGATTTGCA TGCAGGGSTT CATTAAAATC GATGSCAAGS TCCGAACTGA

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30 25 20 5 5 45 6 35 S 50 55 CATTCAGATT GATTTAGAGA CTGGCAAGAT TACTGATTTC ATCAAGTTCC ATTCACCCAG GGTGACTCAT GATGCCCGCA CCATCCGCTA CCCCGATCCC CTCATCAAGG TGAATGATAC CCGTCTGATC TATGACACCA AGGGTCGCTT TGCTGTACAT CGTATTACAC CTGAGGAGGC GRAGCARAGE CTROCEGTRICA CEGAGAGGAA GECLAGETRICT ACTROCEAGGE TRAGECEGTRIG GCAGCTACTG CTTTTCCTCT GTGCCACCCA CTTTGGGGAG CCATTAGAAA AGGTGGCCTC AGCCTCAAGG CACTTCTAGG ACCTGCCTCT TCTCACCAAG, ATGAACTCAC CCAGGTOOTC TCGTCACCTC AGAGGCTCCG CAGACTCCTG CCCAGGCCAG GACTGAGGCA CAAGTACAAG TIGTOCAAAG TGAGAAAGAT CITIGTGGGC ACAAAAGGAA TCCCTCATCT COCCCCCAC AGCCGCCAGA TCCCCGCACC CCAGGGCGCG GTGCTGGTGC AGCGGGAGAA TOTOGGGAAT TOTAGACOCA CAGGOCAGCA GOTAGAATOC CTGGGCCTOC TGGCCCCSGG TATAACCTAC CCTGCTGGAT TCATGGATGT CATCAGCATT GACAAGACGG GAGAGAATTT TITATIGIAT ICIGIAACIA TAGAACIICI ATTIWATICI TITITIGGACI IGCTAAGIIG AAAAAAAAA AAAAAAAAANC TCGGGGGGGG TAAGGGAGGG GGCGCTGGAG CTTCCAACCC GAGGCAATAA AAGAAATGTT GCGTAACTCA ACCAGGGAAC CACGGCAGAA GCGCTGGGGCG GGGCTGAGGG CGCAGGTGCG GGACCTGCCG AACTACAACT GGAACTCCTT CGGCCTGCGC TTCGGCAAGC adeaucence enonceece eceecauaus encessause eceeuseuse essocriste TETTWATGG TITTWAGTIC CATGETGAAG TITTEAGTAT TGACTTATEC CETTGAACAT 3 TIGITYCACIG TITICIGITIGA TICIWACICA IGGIATITITA AFICTICGIT WITITITITIC GAGTTOTTTT ATAGACTOTR ATGATTCAAA AATOTTACAT CTITTOGTAG TOTOTTTCAT ATTACTCACC AGAGAAGATT TTTTTTGTTYT ACCARGTGCC TARGAATGCT AACAGTCTGG TOTTWAGAWA CATTOTTIGA AAAATAATTI GGAGGAATAT TIGATIOTTA TGAACAAGOO INFORMATION FOR SEQ ID NO: 82: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: E CANAGGAGTE AGAGEATTOCG GGGCGGGGGC GGGGGGGCGGG GACGTAGGGC SEQUENCE CHARACTERISTICS: (A) LENGTH: 684 base pairs
(B) TYPE: nucleic acid
(C) STRANDECNESS: double
(D) TOPOLOGY: linear GGGCAGTGAA TOGTTTCTTG 1260 1020 1290 1200 1080 960 900 840 720 660 600 540 480 420 360 240 180 120 360 300 8

> 20 5 . 5 S TITITCTICC GAAAGATICC CCCAACATTA CCATTCCCCA CCTICCCTIG AATTITITIG TTATCAGAGC CCAACTICGA GOGCICIGGG CITTAGCTAC TOTCACCCCA TCATAACTGA CTITICICCC AACCCIGIGG GAATARICAT YCATAICCIA RCTGCAGGCI ARAAGGIGGI GAMCACATAG AMCACCAGGT GATGAGACAA TCCTGGGART CCTGTTTTAC TTTGGSCCAT GCTCTCATTT TGAATTTTTC AAGA GCTTCATGGA TIGATICICT TITTATCTTT CAGATTITCT TITAAAAAATC TITGITTITT (2) INFORMATION FOR SEQ ID NO: 83: Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 2024 base pairs
> (B) TYPE: nucleic acid
> (C) STRANDEDNESS: double
> (D) TOPOLOGY: linear 660 900 540 480 420

25 CTGCAGGAAT TCGGCACAGC TGCGCTGGAG GCTTCATCTT TGCCGCCGCT GCCGTCGCCT SEQUENCE DESCRIPTION: SEQ ID NO: 83:

35 30 GCGCCTCGGG GCTGCGAGGC TGGGGAAGGG GTTGGAGGGG GCTGTTGATC GCCGCGTTTA TOCTOGRAFT GRAGICTORA GCTTTCTTCG TICGTTCGYC GGCGGGTTCG CGCCCTTCTC COSCOSIANCE OGROSOGATO ASGAGGARGA GTOSCTETAT GOCGRIGAAA ATGAAGTTGA AMOGECAGAA GAAGAAAATG CEAGTGETAA TEETECATET GGAATTGAAG ATGAAACTGE AGTTOCOCTC GOOGCGACCA TGTCGGCCGG CGAGGTCGAG CGCCTAGTGT CGGAGCTGAG 360 300 240 180 120

3 8 COATGATGAT GAAGATGATG TTCATGTCAC TATAGGAGAC ATTAAAAACGG GAGCACCACA TOGAACTACA GOGACAAAAG TCAAAGGAGT AGACCTTGAT GCACCTGGAA GCATTAATGG GTATOGGAGT TATOGTACAG CACCTOTAAA TCTTAACATC AAGACAGGG GAAGAGTTTA 600 540 480

TGAAAATGOT OTACCAAAAC CGAAAGTGAC TGAGACCGAA GATGATAGTG ATAGTGACAG

420

TCAAAAACAA AAGAGGATAC GAATGGGACT TGAAGTTATA CCAGTAACCT CTACTACAAA AGTICCACIC TIAGAGGIAG AITIGGATIC TITIGAAGAT AAACCATGGC GIAAACCIGG TOCTGATCTT TCTGATTATT TTAATTATGG GTTTAATGAA GATACCTGGA AAGCTTACTG 660 840 780 720

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GTACAGCAGG GAAGAACTGG AAACTCAGAG AAAGAAACTG CCCTTCCATC GAGITTACIT CICCICCITC TITIGITICAAG ACTOGGCITIC CACCGAGCAG 900

CAGGCGACGG GCAAATGAGA ACAGCAACAT ACAGGTCCTT TCTGAAAGAT CTGCTACTGA AGTAGACAAC AATTITAGCA AACCACCTCC GTTTTTCCCT CCAGGAGCTC CTCCCACTCA GOGGCAATTG ATGITATCGG TCAGACTATA ACTATCAGCC GAGTAGAAGG 1080 1020 960

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×	WO 98/39448 PC	PCT/US98/04493	WO 98/3944§ PC
	305		306
			OCCICATIOC TAAGTOTACT GCTGCCGAAG TTGCCCCAGT TCCATGGGGT TCGTGTCTTT
	CETTOCACCT CCTCCATTTC ITCCACCTCC TCCGACTGTC AGCACTGCTC CACCTCTGAT	1140	OCCATCAACA ANTACTGAGG GATGGGTTTT GGAACAGCTC CATGGGCATG GGGAAGGCAC
v	TECACCACCO GOTTITICCTC CTCCACCAGG CGCTCCACCT CCATCTCTTA TACCAACAAT	1200 5	TGAAACAGAG GACTATAAAA CAICCTICIC TIATICICCA TACIGICTIC TACACCTTIA
•	AGANACTICA CATTOCTICTO OTTATIGATAO TOOTTOTICA OTTOCATTIC CATATOGCAA	1260	AAGCCTGAGA ACTATACAAC CTTTCCCAGA CTCCCAAGAA GAGAAGAT TGGCAAATGG
	TOTTGCCTTT CCCCATCTTC CTGGTTCTGC TCCTTCGTGG CCTAGTCTTG TGGACACCAG		GOCTOCTIGGG COCAGTOCTG CTAGTIGGAA GTTTCTTTGA ATCAGGAAGG CAGGTGAGGT
10	CAAGCAGTOG GACTATTATG CCAGAAGAGA GAAAGACCGA GATAGAGAGA GAGACAGAGA	1380 10	AAGGCCAAA TCACTCTCCT CCATAGCAGG AAGCCATTTG GGCAGCTCCT TTGGTGATTA
	CAGAGAGCGA GACCOTGATC GGGACAGAGA AAGAGAACGC ACCAGAGAGA GAGAGAGGA	1440	CAICTITICCA TATCITITAC ACTIACCACC TICCAGCICT GITTIGCIGT GIATITITICT
:	GOSTGATCAC AGTECTACAC CAAGTGTTT CAACAGGGAT GAAGAAGGAT ACAGATACAG	1500	TACAATAATT TITTICAGCT ATACCICCAG TITTAATCAGG ATGGGTAGAG AGCTGTCCTC
2	GGAATATOCA GAAAGAGOTT ATGAGCOTCA CAGAGCAGT CGAGAAAAAG AAGAACGACA	1560	ATARGECTICG GOOTGOGAAG ATGGAATACT G
	TAGAGAAAGA CGACACAGGG AGAAAGAGGA AACCAGACAT AAGTCTTCTC GAAGTAATAG	1620	
70	TAGACCTICGC CATGAAAGTIG AAGAAGGAGA TAGTVACAGG AGACACAAAC ACAAAAAATC	1680 20	
	TANAGANGC ANAGANGGAN ANGANGCGGG CAGTONGCCT GCCCCTGANC AGGAGAGCAC	1740	(2) INFORMATION FOR SEQ ID NO: 85:
;	CGAAGCTACA CCTGCAGAAT AGGCATGGTT TTGGCCTTTT GTGTATATTA GTACCAGAAG	1800	(1) SEQUENCE CHARACTERISTICS: (A) LENTH: 825 base pairs
22	tagatactat aaatctigit attitictgg ataatgitta agaaaittac citaaatctt	1860	
	GITCIGITIG ITAGIAIGAA AAGITAACIT TITITCCAAA ATAAAAGAGI GAATITITCA	1920	(b) TOPOLOGY: linear
30	TOTTAGOTTA AAARCTITG TCTTGTACTA TTTCAAAAT AAAAGACAG CAATGACTTT	1980 30	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 85:
	atatccaaaa aaaaaaaaa aaaaaaaaa aaaaaaggc ggcc	2024	COGGOCCOGC GGOSTCTTCA GGSTACCGGG CTGGTTACAG CAGCTCTACC CCTCACGACG
		S C	CAAACATGGC AGGGCAGAAG GACCAGCAGA AAGATGCCGA GGCGGAAAGGG CTGAGGGGCA
32			CONCECTOR OCCOMMENTS AFFECTIONS OFFICIALISES GANGTOSCITO GAGGGGGGCC
	(2) INFORMATION FOR SEQ ID NO: 84:		GEGEGACCAT COGCOCOTIGG AGCACCTTCG TGGACCAGCA GCGCTTCTCA CGGCCCCGCA
40	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 911 base pairs	40	ACCIGGRADA GCTGTGCCAG CECCTCGTAC GCACGTGGA GTACTACCAG AGCAACTATG
!	(B) TYPE: nucleic acid , (C) STRANDEDNESS: double		TOTICOSTOTIS CCTGGGCCTC ATCCTGTACT GTGTGGTGAC GTCCCCTATG TTGCTGGTGG
	(p) TOPOLOGY: linear	•	CICTROCTOT CITITICOSC GCCTOTIACA TICTCIANCT GCGCACCITG GAGTCCAAGC
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:	45	TIGHCCICTT TGGCCGAGAG GTGAGCCCAG GGCATCAGTA TGCTCTGGCT GGAGGCATCT
	CECCECEMATA GCCGGAGGGG GATCTGAGCT GGCAGGATGA ATGTGGGGGGT GGCACACAG	09	CCTTCCCCTT CTTCTGGCTG GCTGGGCG GCTCGGCCGT CTTCTGGGTG CTGGGAGCCA
S	GAAGTAAACC CCAACACCCG AGTGATGAAT AGCGGAGGCA TCTGGCTGGC CTACATCATC	120 50	COCHAMINAT CATOGGCTICC CACGCTGCCT TCCACCAGAT TGAGGCTGTG GACGGGGAGG
2	THOGRAGOAT TOCHOCATAT GGTTCTACTC AGCATCCCCT TCTTCAGCAT TCCHGTTGTC	180	ACCITICAGAT GRANCECOTIC TEAGOTISTET TETGGGACET GEOGGEETEC CGGGCCAGET
	TOGACCCTGA CCAACGTCAT CCATAACCTG GCTACGTATG TCTTCCTTCA TACGGTGAAA	240	GOCCOACCC TGCCATGCC TGTCCTGCAC GGCTCTGCTG CTCGGGGCCCA CAGGGCGTC
55	GOGACACCCT TTGAGACTCC TGACCAAGGA AAGGCTGGGC TACTGACACA CTGGGAGGAA	300 55	CCATCACAAG CCCGGGGAGG GATCCCGCCT TTGAAAATAA AGCTGTTATG GGTGTCATTC
	ATEGRATATE GOCTOCAGTY TACCTOTTOC COCAAGITOC TCAGCATCTC TCCTATTGTG	360	ACSAAAAAA AAAAAAAGG GGGCCCCTC TAGGGGTCAA AGTTA
9	CTCTATCTCC TGGCCAGCTT CTATACCAAG TATGATGCTG CGCACTTCCT CATCAACACA	420 60	
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TOGRAGARCE CARACTRAGA CAGTOCTCCT GGTGCCCT

INFORMATION FOR SEQ ID NO: 87: SEQUENCE CHARACTERISTICS:

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AACAGGGGA GCCCCAAGGG CTCGGTGCTA TTTGTAACGG GATTAAAATT TGTAGCCAG

CAGACACTAC ATGGGTAGCT CAGGGGAGGA GGTGGGGGGTC CAGGAGGGGG ATCCCTCTCC

1380 1320 1260 1200

1440

COMPAGENTA GEOGRAPHIC CAGAGGGACC CIGGGCCGIG IGCCAGCTCC ACCIDEACAT CIBECACAGE INSCRETINGS COCACCECAE GAAGGGCCTG GOCCOCTICEA COCGOGGIGEC COTGGCCAGT GTGCTCATCT GGAGAGGCTG

1080

960 900

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ACAGCTOCAC

GIGCAGGGCC

TOCTOCACGA AGCAGGGTCC ACAGACGACC GGATTGTGGT

1140 1080 1020

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TOCTTOTION COGNOTICOG GEOTOTICAG TICTOGCICA TECTOTICOG GITTETICOG OCTOGRAGE CETACETCAT TOGECTGATE TETGACCOCC TOGGCCGGAA CTGGCCCCCC CCTACCCGAC GCTCCACCGC CGAGGCCTTC CAGATCGTGC TGTCCCACCT GCTGGGTGAT

> 960 900 840 780 720 660 600

GCGCACTTCC TGGGCACCGC CATCTTCATT GAGGCCGACC GCCGGCGGGC

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GAGACCCTCC TOTCCATGAA CTOGGCCATC GTOGCCGACA TTCTGCTGTA CGTGGTGATC CIGICCCITG CCIGCGCCC TGGIAGCAIC GIGGCCACIT ANAITITICAT CITCATIGGA CCCCGGGCIG AICCCCIGGT CIGIOCCACT GGCCICCIGG GCICIGCACC CIICCICTIC

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780 720

3 6 35 3 50 GOGACOGTIT CTOGCTCTCA GGCTCTGAGA AGCTGCAGTT TATGAGTGGC TCTGTGTGTG CATGIAAAAG GAIGAAAIGI GACIICIGGI GIITITITIAI TICIAIGGAG GGACITICIG GCTGGCTCAC TGCTCTGGCT TCATTTTCCA GAGCTGCCTG CTGCAGTCAC ACTTAGGTCA CAGGAAGGOT GTAAGGAGAG GATGGATCCT GATACATGGA TICAGGATCA TIAGGGICCT TOTGGGGAAG TOGGATOGCA GCATOGCAGG OCTITTOGAAA ATGAGAGGTG AGAGTKTKTC CTGCCACCTA CTGGAGAAGC CATAAGCTGC AGCTTTAGGA AAAGGGAACC CGGGGCAGAG TOCCCACTIT TITTCTOTOT CCTGACAAAG AAACACAGAG TAACTTGATT GCCCTGTGAC TTCACTTOCA GCCOTGATGG AMAOTTNOCA TOGAAGCTGA GACTCTCACT GACAGTGAAA GGGATGTCCT TTGATGGCAT CAAGACTTTA GCTTCTGGTG CGCTGTGTCC CAGCTCTGAT AGGTTOTTOC CCAAAACACT GAAAAAAAACT GCCCTGGCCC TGAACCAAAT ACCTTGAACC CTOGCCAGTT GCATTTCCCC TGCAGGCTTG AGCCCAAGCC AGAGCCTTGA AAAGGTATTC COCTOMANTO AMEMICANTICO CTECTTTICCT GCCAMGGATC CTIGITAGGGT NCCCCCAGCT TETTETETEA CITITETECT TITIGECGATT AGITGACGIG ACAGAGATOT GAATGOGGCA GICTOGGACA CIGOCCTICC IGCTIACCIG CICITICCII CCICCIIGGI CGGAGGAGG TOCTTIGGAC TGATCACCCT GCCAGTCTTT TGTCTTGGGC AATCTATACT TTTNCTCAGA CTCACTOTCT GTTGTGAGGA TACGCTGTAG CCCACTCATT AAGTACATTC TCCTAATAAA CTCGTAAACT CCATACCCTG ACCCCCTTGT TTTGGATATA CCCAGGTAGA ACAACTCTCT OCCUTTAGET TUANGAATAC AGGATUACUT GTACCUAAGC CUTTAGUUCA AGUTUTGUTT TTACCTTATG CCCTCACTTC CTGAGTTAAC CTCCCAAATA CAGGATTCAC CTGTACCCAA GOTTECCAAG GECTACTGAA GOGACTTAAC ATACTETTAA TEGETITEET CTETETTETT (1) SEQUENCE CHARACTERISTICS: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 86: g 0 9 TOPOLOGY: linear STRANDEDNESS: double TYPE: nucleic acid LENGTH: 1238 base pairs 1020

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TIGCCACCCC IGAACCCCAC CICGIGGIGG GCAGAICIGA GGGCICIGGC AAGAAAICCI

AGITICGICC IGICIICCCI GGGCTICACI GCIGIGGCCI TIGICACGG CICCCIGGCI

480 420 360 300

CTGCTGCTGT TCCTGGTAGT GCGGGAGCCG CCAAGGGGAG CCGTGGAGCG CCACTCAGAT

CTOTOGOCTE COGCATTECT OCTOCOTTEC COCOTOGICE TTOGGGAGAC CECACECTOC

ACCOGAGICC TODGIGIOGG CCITOGGIGIG GAGATCAGCC OCCGGCICCG CCACICCAAC CTICCCOGAG ACTOCIOCIC TICCICIGAC AGICICAICI TIOGACICAI CACCIGCCIG 15

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CTOCTCCTCC TGACCCGGGG CCTGGTGGGG GTCGGGGAGG CCAGTTATTC CACCATCGCG

ATTOCCTTCT OGTCCCTGGT GACACIGGGG TCATCCTTCA TCCCCGGAGA GCATTICTGG

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SEQUENCE DESCRIPTION: SEQ ID NO:

(A) LENGTH: 1460 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

TACTITICCCA TICCGGIGGG CAGIGGICIG GGCTACATIG CAGGCTCCAA AGIGAAGGAT CCCACTOTCA TIGCOGACOT CITIGIOGOC GACCAGOGO ACCGGATGOT CAGCATOTIC

240

9 120

ANGOCTOGAG ACTOGCACTO GGCTCTGAGG GTGACACCOG GTCTAGGAGT GGTGGCCGTT

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(2) INFORMATION FOR SEQ ID NO: 86:

PCT/US98/04493

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PCT/US98/04493

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\$	WO 98/39448 PC	PCT/US98/04493	WO 98/39448
	309		310
	(2) INFORMATION FOR SEQ ID NO: 88:		OB ON AT AND INVESTIGATION (17)
	(1) SEQUENCE CHARACTERISTICS:		
ν.			5 SAQUENCE CHANCLERISIS.  (A) LENGTH: 1186 base pairs  (B) TYPE: nucleic acid  (C) STRANDEINESS: double
;	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 88:		(D) TOPOLOGY: linear
<b>9</b>	CAGGIGGAAA GTGGGAAGTG TGAGTCCTCA GTCTTGGGCT ATTCGGCCAC GTGCCTGCGG	09	10 (xi) sequence description: seq id no: 89:
	GACATGGGAC GCTGGAGGGT CAGCAGCGTG GAGTCCTGGC CTTTTGCGTC CACGGGTGGG	120	GGCACGAGCC GGCAAGCCGA GCTAGGGTGA AAACTGGGGG CGCACCAGGA TGTNNGACAG
15	AANTIGGOCA TIGCCAGGGC GGGAACTIGGG ACTCAGGCTG CCCCCCGGC GTITCTCATC '		AAAAGAGAA GATGAGACTC TGTTCATTCA CTTTTCCTAG GCCCATCCTG TGGTCATCTT 15
	COTCCACCOS ANTICOTOSSIC GOTCOCACTG GOCCTGATOT ACTITICCTGA COTCTGACCO	240	TOCCOCTOCC ATCATACCTC CTCCTTCCTG GAGCTCTGC CGGCTTGGCT GTAATGGTGG
ć	STATISTICIC CAGNITAAAG STACGACATT TGGAGGCCCC AGCGAGAAAC STCACCGGGA	300	
9	GANACITICAC COGOCCIAGAG COGICCOCOT GTOTOCTICOC COGGNAGGAC AGOCAGOTTIG	360	20 AGANGGOAG AGGAAAAAGG CAGAGATGGC CAGGAGAGGG GTGCAGGACA AACCAGAGAG
	THGGGGGGAG TCCCACCTGA AAAAAAATT TCCAGGTCCC CAAAGGGTGA CCGTCTTCCG	420	OTTOGOTICAG GCGAAAAGGG TGGGGAGAAA GAGGGGTTGCA GGCCCTTGCAG GCCGGTTTAGC
25	GAGACAGOGG ATCGACTACC ATGTGGGTGC CCACAAAAAT TYCACCTYTG AGTCCTCAAC	480	Chechectise secretoes seccritises attendente scalacasse taccaseere 25
	TYCTISACCCC GGGGTCAGTT CCAGAGAGA GGACTCCCTC CTGCTTGGAA GAGACCTCAC	540	CTGGTGTGTA TCATAGGATT TGTTCACATA GTGTTATGCA TGATCTTCGT AAGGTTAAGA
Ş	ACCOTCATCA CGATGCCAAC GGCTCTGAAG GTGGATGGCA TTCCTGCGTG GATTCATCAC	009	
Sc.	TCCCGCATCA AAAAGGCCAA CRGAGCCCAA CTAGAAACAT GGGTCCCCAG GGCTGGGTCA	099	3U TOTATOTANA TTATGGCAGG AGNANTATA GCACTGAGGG CCCTGCTGCC CTGCTGGAGC
	GOCCCCTIAN ANCIGENCET ANGINGOSTG ANGCENTING ATTANTICET TITICITANT	720	AAGCAAAACT AAGCCFTTTG GTTTGGGTAF TATGTTTCGT TYTGTTAFTT GTTTGFTTTT
35	TIGITADACA ATGCATAGCT TCTGTCAACT TATGTATCTT AAGACTCAAT ATAACCCCCT	780	GIGGCTIGIC TTAIGTCGTG ATACACAAG TGCCAGTCGG ATTGCTCTGT ATTACAGAAT 35
	TOTTATAACT CAGGGAATCA ATGATTTGAT TCCCCAAAAA CACAAGTGGG GAATGTAGTG	840	AGIOTITITIA ATTCARCAAT GITCHAGITA ATGICTAACCT CAGCACCTCC TCTTAGCCTA
Ş	TECANCETES TITITACTAN CECTSTITIT AGACTATICE TITICETTINA TEACTEMENE	006	
<b>4</b> . Ō.	TIGITICCAC CIGAATIGAC TCTCCCTIAG CIAAGAGCGC CAGAIGGACT CCAICTIGGC	096	40 caarciety cicietytes crociocosa ecicicados tysocorcio carciocore
	TOTITICANCT GOCKGCGGT TOCTYCAAGG ACTTAACTTG TGCAAGCTGA CTCCCAGCAC	1020	TOCOGCOCTO COCTOCTOCO TOTOGOCTOCO COTOCOATTA CAGACACTO CATTOTOCTOCA
45	ATCCAMBAT GCANTTAACT GATAAGATAC TGTGGCAAGC TATATCCGCA GTTCCCCAGGA	1080	септетета выстастве твосестое семессмет тососмент втоетносов
	ATTICGLICIAA TIGAITIACAC CCHAAAGOCC CGCOTCIAIC ACCTIGIAAT AAICTIAAAG	1140	CTCCTTATCT GTTCTAGTTC CGAAGCAGTT TCACTCGAAG TTGTGCAGTC CTGGTTGCAG
S	CCCCTGGAC TGGAACTATT AACGTTCCTG TAACCATTTA TCCTTTTAAC TTTTTTGCCT	1200	
ક	ACTITATITIC TOTAMANTIC TITTANCING ACCCCCCTC TCCTTICIAN ACCAMBINE	1260	JU CIGACAGGAT TIPAAAAAAAA AAAAAGGAAA AAAAAAA AAAAAA
	ANANGCANAT, CTNGCCCCTT CTTCNGGCCG AGAGANTTIC GAGGGTTNGC CGTCTCTTGG	1320	
55	CCACCAGCTA AATAAACGGA TTCTTCATGT GTAAAAAAA AAAAAAAAA CTCGGAGGGG	1380	55 (2) INFORMATION FOR SEQ ID NO: 90:
	מספכתמסתות מכבאת	1395	(i) SEQUENCE CHARACTERISTICS:
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900 960 1020 1080

780 840

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1140 1200 1080 1020 540 1380 1320 1260 960 840 780 720 660 600 480 420 360 300 240 180 120 1680 1620 1560 1500 1440 900 25 20 8 45 8 ઝ ဗ 15 5 8 55 TITTAAAATG TGACTCTCCC AGAGAAGAAG CCGCTGGCTG TATGAAACTT GACGGCGCTT TOTTTECTION THTCTOTICCT TECCHTTTTA CAGGACTECE GGAAGGECAE TEATGGECAT TOCCCTTTTT CCCACCGATT COGGGCNTGG TGAAGGTGGG AGATGTGAAC TCCAATTAAG (2) INFORMATION FOR SEQ ID NO: 91: AACCCGGGG GGGTTTCCCC C GAGGGOTTAT TITNOGCTTG GGCACTGGGC CCTTCGTTTT TACAACGTCG TGANGGGGG AAAAAAAAA AAAAAAAAA AAAAAATTC CTGCGGCCCG CANGCTTTTT CCCTTTGGGT CTTGTTCATC CTCCAGATGT AGCTATTGAT GTACACTTCG CAACGGAGTG TCTGAAATTG CTCACCCTCA CAATITATITI CCTCCTCCCG TGCCAGCCCT TCTTTTGTGT GACGITITIC IGCICICITC IGGCCCICCA IGGAGCCAIG GGCCICGGCC ICGGCGGCTC GCCAGGAGCT TICICAGAAA CAGICATAAA CGAICICIIG AGICICITIC TIGICCICCC TACCACCOCT TOGAGTCTCC CGAGGACACA AACAGGCAGA GAGGGACGTG TAGGGAGAGT AAATGACTIG ACCTIGCCAT CIGIGITICAA GGICACGGIT IGCIGIGGGG TICCIGGGAG GGACTOGAGA GAGGTGAAGA ATTITIGCAGG TOOGAGAITT GGATTTGAAIT GTGGACTTOT TIGIAAGGIG CCACCCCCAA ACTITIAAGGI AGCTGAGCTT TCTTATTCCA CCCTTTCTGG TGTCTATAGG AATGCATGAG AAGACCCTGG AGCTTACTCA CCCCGGAGTC TTTTCTTTCT CTTGCTCCAA GAAGAGCCCT GTTGGTGCTT AAAAACTCGA GGGGGGCCCG GT 2 INFORMATION FOR SEQ ID NO: 92: Ξ (x1) SEQUENCE DESCRIPTION: SEQ ID NO: £ SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: (A) LENGTH: 862 base pairs
(B) TYPE: nucleic acid
(C) STRANDECNESS: double
(D) TOPOLOGY: linear (A) LENOTH: 696 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear AGCTAAACCA ATTITITAAAA GATTCAATGG 91: 1800 1740 1821 540 420 360 8 240 180 120 480 862 840 780 720 660 600 60

30

GCAGTGAGGA TGTTTCCAGA

AGCTACTTAC CTACATGTGA

ACTITITIGGA ACAGTAATGG CAGCTICTAG TACAGCCATT

TIGIAGAGAA AAICCCCITT

TACTAAAACA AATGACACTC TAAGAAAGTT TOGGAGCCCC

ATGCTGAGAA CCATTTCTGT AIGIGCCATT TICTITICCTI

3

ACAGTTTCAT ATGAGAAAAA TIXAGAATAA CTATAAAATT GTTAAAATAT CCAATAATGG

ATAATGATGG CCAGAAGATT TAACATACAA AGTAATTCTC AATGTAAAGC TATTCAGGTC

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AATGCCCTGT

AACCCACCCT GACCTTCCAC ATCATCTTCA AAAAGCAGTT

rererence TICCAGGTIG

45

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CATTOTICO TOCCTOCCAT CAGGICAAAT CAGGAGGOTG CAGTGAATGC CTGTTCTTTG

AATTAGAGCA AAACAAATCC TCTACAAATC CAAGGCAGGA AAAGTGGTGG CAGAGTGACT

TITCATGCCC TATAAGCAGG TACCTTAGTA GGGCAGATAT AGGAAAAACA ANGANTAAGT ATCATGTGAT TATTTTAGCT TTACAAAAAA AAAGTTGAAT CCATGATICT CCTATAAGGT AACTCTTTAG TCCTCCATTT AGCACATTTT

ANTOTOTAGO AGTIGITICOT GIAACIOITI AAAACITOGO TATAGGCIGI TIAGCACAGI

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TCARGCTING GITTOCTAAT TITTITCACAT ACCTITITICT ATCACAGICT GITOCTITIG ACAGATTAAA GATACAGTTA CGTAAACAGC AAAGTAATTT TATAGTGCTT CATCCATTTA

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TACACATTIC ICATATIGGG GIICGACAGG TAAACACAAA CIGCIATTIC AGIAGAAAA

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CTARTIAGAA GGGGAAGTIA GCCACAGAAA ATCAACTIAT CTATAATTAC AAAATTCICT CAGCAGNAGO AACCCCACCA GCCTAAGTCC AGCAGAGGAC CTCCCACCCA ATGTCTTGTT GTGGCTATCC CTACAAGGCC CACCCAAGGG ATGCCCAAAG CCCAACCTTC TCCAGGGCTG AGICAAATCA CTACTICCAT TOCTACTITA GATCAGCCAA AGIGGIGACT GCTGCAGIGI CACCCAGACA CAAGCCCCTT TCCCAGGTCA AACCACAGGC CGATGCATCT CCAGTTTGAC

TGACTCACCT TAAAGTTCCT ATTGACATCT ACTOCTTTTA AACCTATTTG AAAACTCTGA

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TGATTCAACA CAGCTOCCCA CACAAAGCCA GTOGTAATAC ATCTGTTTAC CTTTCCCTAT

CTCCATCCTA GAAGCCAGCC CTAGGMAGCT GCAGTTACTC CCTGTGACTC AGCAGCAGGC

GGAGTITICAC ACACAATGAC AGGCTGCTGG GGGACATTGC AGGACCCCTT TTCCTYTCCT

ACAATCCCTC CCAGAAGTCT CCCAACACTA GTGCTGACCA GAGGTGGGGC TCTCAGGCTA KATCCOTCCT CACTOCGCTC AAGATGGCCT CAGCAGACAC CAGTTACCCA GCTGAAAGTC

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SEQUENCE DESCRIPTION: SEQ ID NO: 90:

(D) TOPOLOGY: linear

AAAACATOCT TICAGGGCGT CCCCTATGTA TICGGGGGGC CCACGGACAC TCAGGCTGGA

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 92:		CTGAAGTCTA TATCGGCATC GGGAAGCCTG CAGAAGCCAC AGCCTGTACC CAAGAAGCTG
CTGAGGGAG TGAAGTGGAC TCTGAGGGCT ACCGCTACCG CCACTGCTGC GGCAGGGGCG	09	CCAACCTETT CCCAATGTCC CACAATGTCC TCTACATGCG CGGCCAGATT GCTGAGCTCC
TOGAGGGCAG AGGGCGGG AGGCCGCAGT TGCAAACATG GCTCAGAGCA GAGACGGCGG	120	
AAACCCOTTC GCCGAGCCCA GCGAGCTTGA CAACCCCTTT CAGGACCCAG CTGTGATCCA		CACTIGAAGA GCAITCCAGG ACTIGGCCCT GAITCCTTCAC CAGTIAGGCC GYTACAGTYT
GCACCGACCC AGCCGGCAGT ATGCCACGCT TGACGTCTAC AACCCTTTTG AGACCGGGA	240 10	
GOCACCACCA GCCTATIGAGC CTCCAGCCCC TGCCCCATTG CCTCCACCCT CAGCTCCTC	300	
CTTGCAGCCC TCGAGAAAGC TCAGCCCAC AGAACCTAAG AACTATGGCT CATACAGCAC	360	
TEAGSCETEA GETGCAGEAG CEACAGETGA GCTGCTGAAG AAACAGGAGG AGCTCAAGGG	15	
GAAGGCAAAG GAGTTGGACC GAAGGAGGA GAGCTGCAGC ATGCTGCCCT GAGRGGCACA	480	АССАВЗЕСТ ВТЕССАТОВС СОСААБЗВБА ТВАЯТСТВСС ВСАСТВАВОС САВОВАСВАБ
GCTACTOGAC AGAACAATTG GCCCCTCTA CCTTCTTTTT GTCCAGTTCA GCCCTGCTTT	540	
TICCAGGACA TCTCCATGGA GATCCCCCAA GAATTICAGA AGACTISTATC CACCATGTAC	009	COTTICTION STITITITIE CCAGCCCAAT GGIAGITICE GAACCIAITG ACATIGITICA
TACCICIGGA TGTGCAGCAC GSTGGNTCTT CTCCTGAAYT TCHTGGSCTG CCTGGCCAGT	099	AAATGGAICA TGTGCCATAF FTTGTFAGTT GACATCTGAG TTTTCAGTAA AATGAFTANG
TCTGTGTGGA AACCAACAAT GGGGAGGCTT TGGGTT	696 2.5	
		TATTITIATE AGAGCISTA AGANAGANE TGTOCTTITC TCCCCACCAE CCCTCCTGCC
(2) INFORMATION FOR SEQ ID NO: 93:	30	CCACTITIGGC CCAGAAACCA AAIGTGAACT TCCTGTCTCC CACCTCAGCA CTAGTCCATG
(1) SEQUENCE CHARACTERISTICS:		CCAGGACACC ACCTGACAAT TYCTYGGYYY TACYGYCAAT AAYYGYACCA TGYGAYCAAY
(A) LENGTH: 1886 base pairs (B) TYPE: nucleic acid	V C	TACTISTICS ACTINGMEN ASSOCTIONST COGGONIAN THAINTITTA CCANINING
(C) STRANDEDNESS: double (D) TOPOLOGY: linear	cc	CCTGITACAA GAGAAGGAAA TAIGAGITAT TIAAGITTAA CITTITIATG TGAAITCAGA
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:		GTTTATTTAT CCAGGGAAAT ATGTACAAAG AAGCTTCAAA TOGAATATTT ACCGACATTC
CAGGCCACTG ACOCTTCTTT GCGAGGGATG CAGGAGGTCC TACAGAGAAA GGCGCTTCTT	60	CTIMINCATG ACAGACACTT GOCTACATGG GAAGATGATG TIAMINATAA AMIGATTITI
GCATITICAGA GOGOCCACAG CCTOTCACCC ACAGATCACC AAGCAGCTTT CTACCTOGCT	120	ааатосалла Лалалалла Лалали
CTGCAGCTTG CCATCTCCAG ACAGATCCCA GAGGCTCTGG GGTATGTCCG CCAAGCTCTT	180	
CACCITICAAG GIGACGAIGC CAACITCCTIG CCCTCCTGCT GICAGALAG	240	(2) INPORMATION FOR SED ID NO: 94:
ANGCATTACC ATGACGCTCT GARCATCATC GACATGGCCC TGAGTGAATA CCCAGAAAAT	300	(i) SEOTEMER CHARACTERISTICS
TICATACTAC TOTTTICCAA AOTGAAGTTG CAGTCACTCT GCGAAGGCC GGACGARGCA	360 50	į
CTGCTGACTT GTAAGCACAT GCTGCAGATA TGGAAATCCT GCTACAACCT CACCAACCC	420	
AGIGATTETE GACCTOGGAG CACCOCTTA GATAGAACCA TTGCTGACAG ACGACAGCTT	480 55	(xi)·Seque
ANTACANTTA CTTTGCCAGA CTTCAGGGAT CCCGAGACAG GCTCCGTCCA TGCCACATCG	540	CTCAGCTACC GTATACAGTA GOACATAACC CCATTTCACA TGCACTACAC TGAGACTTGC
GTAGCAGCCT CAAGAGTGGA GCAGGCACTG TCGGAAGTGG CTTCGTCTCT GCAGAGCATG	009	CTCCTCTCCC CCCACATIGA AGAIGITCIT TITICAIAAC TAIATACTAT TCCATIGCAT
CCCCTANGCA GGGCCGGCTG CACCCTIGGA TGACGCTGGC ACAGATCTGG CTCCATGCAG	09 099	

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GAATATICIG TAATITATIT AAICCCCIAI GGAIIGATAA TIAGGIICAT TATAGATAGA

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GOGGAACCIA TICCCIGIGG CITAGGIGAG CAIGIGACCA GOCCIGOCCI CCIGAGICCC TITTIATING TICAGCANCI CCTICCCCAT CINCIGGIAA CACAACCITI ATTIATING TTACIGIOGA TIGITIGIAT CCCTTACCIG CITICIATIG GGITATGIGI GGATATATIG TGATTATAAA AAAAAATGGT GAGATTGGGG TTATTTTCAT GTTTATTGGC CATTTATAGT TACACTIGAT ACTITITATIC TOTTIGGCCGA AAAAGAACCT TITCITATIT TGCATTITCCC TICATAACIG TATTITICACC AAGIGTAIGG AGAAIGITCA TITICCCCATA TAACCATACC

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GIGGGGSCIG

COTOTTOCTO TOCTATOTOT GUALIGOGIC ARGOTOTOCO TOCOGGGCO TGGGTTTCTA

660 600 540 480 420 360 300 240 180

GLAGIGGGCT TECTOGCIGT KITECTETOTO CIGCIGGCAG GCGGCCTIGGG

SCICIOSSEA GOSCCAGSE CITACICAIC CICTISCITA TAGCCATOSE TOTOTICCCI

ACAGCTTCCT AGCCACAGTG ATAAAAGAAT GGGTATATAA CTTAAGCCAG GCTAAGGAAA

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GICCCITICI

GATICEAGETG TECCTAAAGE CICCCCTACT CCGGACTITA AAGITTTGTG AGCCAATAAA CGGATGCTAG CAGAAATGGA AAGAGAACTA AGTTCTGATG TCATTTTTCT GGAGGCCCTA CCAAGGACCA TOTAAGCCTG AATTTOTOCC ATGTGGAGAG AGTCTGTCTG AGGAGAAACT GCCCTTAACA GAACTICIGC IGGAACTACT GGAAAGAAGG CTITAIGGAG AICCCAGGAA

> 1560 1500 1440

TGITTAAGAT AATTGAATTG AGTTTCTCTT CTGATTAATA TAGGTTATTT

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ATATAAGITA ATAAAATTAG CATGGCCTTC CATG

GTATTTTCTT ATTGATTTGT AGAAAACCTT TGTAATTTTA AATTCTAGAC TTTATGCACT

1740

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CTCCACTTCC CAACCCAGAA CTTGGAAAGA CATTAGCACA ACTTACGCAT TGGGGAATTG

1560 1500 1440 1380 1320 1260 1200 1140 1080

CCTCGCTGTG AACAGGACTG GACGGTTTGCG CACAAACAAA CGCTGCCACC

acerececte

ACAMATCACT CCCATGATGA GACCCTGGAG

1680 1620

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ACTOGRACTE CARCECCAGA GGTECAGGAG TGATETETGA GTGACTEAAG GGACATOGIT AAGGAGCITC CACTCAGCCC ACCATAGIGA GIGGGCCGCC

ACAAAGACAA GGCTTGACTG CTTCAAAGCT TCCCTGGACC

ATGAAACCTA GCAAAGAACT TACGGCAACA

AACGAGGACA TTAAAAGAGC GAGCACCTCA

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ACACCAATTC CIGCITTAAT TAATGGATCT GAGCAAATCT TCCTCTAGCT

TGATTGCTGT CATGGGGCCA GACTTCCAGG

CTGATTTGCC AAATGCCAAA

TCAGGAGGGT

6

GICTGICCIT

CAAAGAGGCC GAGGGGCAGC AAGGGCAGMC AGGGCACCTG TGACTTCTTA GTACAAGATT

CAGGACTICC AAGGCTCCCA AAGACTCCCT AAACCATGCA GCTCATTGTC

1020

960 900 840 780 720

TAACGCTGAT CICCAGCTCC AGCGATGGAA CCCACTACAG AGGAGGTGGG GCCCCTGTGT GOGTTCAGTT CCAACCATGG TCAGAGGTGG CACATCTGCT CAGCCATCTC ATTITACAGC CTGAGGGCTG AGAGGGCTGA GAGCAAGCTT GAGAGCTGCT AAAGGCTTAC GTGATTGCAA

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INFORMATION FOR SEQ ID NO: 95:

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ACATCCTATT CAGATGTCTT

AMGGTGGATT TTTGGATCAA

GAAAATGCCA TTATTTATTC CCTTAATTTT TTTCCTCTCG CTATTACATT GCCAAAGTAA

TGTGCATGTG TGTGAATATT TCTTTAGTCT GGAGTCCAGT AGGGITTIGIT CICIGICCAC CITCAGICIT CCCAAAGGCC

960 900 840 780 720 660 600

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CCTCACCCTC TITIGCCCCCC AGCCTCTCCT CCTAGCCCAG TOCAACAKTO ATGAGAGAGAG consecrate consecrate econsecon seconsecrate econstants esteconsec CTGCCTGTGG AATGAGGACA CCAGCACCCT ACAGTGTCAC CAGTTCCCTG AGCTGGAAGC TOTOTOGOAG GOTOGOAACO TOAOTGACOT GOOCAACOTG AGAATOGGOT TOTATAACTT GCTOTTCATO AGCATCATAG TCCTCGTGAT TOTGGTCATC TGCCTGATGT TATACGCTCT

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CTCAAAAAAA GATAACAGGT

CTITAAATTT CTTTAATGGT TGAATATGAT TAAATACTAT

CATAAAAACA CITTITACAT AAATAGGATC TCATATICTG TAGCTITITA AAATTITGGT

ACAGITIMAC ATACATOCTO TAGTOACCIT TOOGTACGAA TATACATACA

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GTATAAAAAA CACAGTTCTT

TOCACTTACC

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GAATATTAAT GAGGAGGAAC ATCTTTTCAT GTTTCTTGGC CATTTGCATT TCCTATTATG 

> 480 420 360 300 240 180

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SEQUENCE DESCRIPTION: SEQ ID NO: 95:

GOCACGAGCG AAGGCAAGGG GOCACCAGCT CAGGACTGCA TCTGCCTGCC ATTTCCCTTC

AACCCIGARY TOCTOCTICE CONTINUANG AGRICATOCIG TOGTCICIGG ACCICCITIC

TOTOCOCOG CAGAGGGCAG TAGAGATGGC CGGCCCAAGG CCTCRGTGGC GCGACCAGCT

CACTOCTOCT TTOTOGRAGIC TGACATTAGA AAGOCAGOGA GAAGGAAGAT TCAAACAACG

120

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SEQUENCE CHARACTERISTICS

(B) TYPE: nucleic acid (A) LENGTH: 2503 base pairs STRANDEDNESS: double TOPOLOGY: linear

GCCCATTITIC CITITITIAA TIAIGAAAGI CIAAIGACIA CCIICICATI

TGAATAGAGA GACCCITTTC TCCAATGCTA CCAATCACAT

S

AATAACTAGA AATITATIGG ATCAGGTITC ACATTIGCAG TITIGAAAAC TACTACCAAA

agigiaatta acaticcigt acatgiatit tectactigt giesstatit cigiaggate

AAGATITICAC CAATITIACAA CTCCATCATT AGTAAGAATG CCTGTTTGCC TATAGTCTGC

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			WO 98/39448	PCT/US
	317		318	
	TOTOTRATITI CTACTACTIC TOTANICIA AACCIGIANG GCAGIGATIT ATTCATARA	1620	GANCANCECE OCEOCENGES OCCENTRACE NASTECECES CARCECTORS COSCETTECE	<del>4</del>
			GOCCOGGOC GETECCHANG CAAAGAAGC CECTGAGAAG TECACCTAGT TEACAGGATA	ν,
Ś		5	AAATCCCACA GCAGAACTCG GAGTCAGCAA TGGCTAAGCC CCAGGTGGTT GTAGCTCCTG	
			TAITAANSIC TAAGCIGICT GIGAANGCCC CIGAANTITA CCCTICAGGT TAITICTICCA	. 9
9			GITACACAGA AȚECTATGAG GATGOTISTS AGGATIATEC TACTETATEA GAATATET	- 2
2	TATACACTOT GEOCTICAACC TOCCAGACAG GEOLGAGAAC TOTGGGCAGC TOSTITUCTT		ACCATITITI GAATCATCIT ACACACCAC CTCCCAGITT TGAAACTGAA ATTGAACAGT	
	ICTAGGCTGG CTGGAGGT GGAAGCTCAT TGATAGACTC ATGATGGAAA CTATTTTTGA	1920	TICCAGAGAC CCTGAATGGT TGTGTTACAA CAGATGATGC TTTGCAAGAA CTTGTGGAAC	
<u>~</u>	AACAGGCTTC CTCCTTCAGG AGAGATCATG CGGACTAAAC TGTAGCAATT CCAGTGCACC	1980 15	TCATCTATCA ACAGGCCACA TCTATCCCAA ATTTCTCTTA TATGGGAGCT CGCCTGTGTA	8
3	TOGCHOTGAT CCTTTTCTTT GCAAAGTACT GTCTCTTTGG TTCCAGTAAG TTGGACCACC	2040	ATACCION CONTONICIO ACANTINOCO CACACANOS CAACITOCOS CAATITOCIAC	
	ACATGACATY ATTITECETG GAACETGGTE ACTGACTAAE ACAGACAATT GGGACTECAG	2100	THE STATE OF THE STATES AND STATE	5
20	AGCCTCAAGA GCCAGGAGAG GGCACAGTAC ATACAGAGG AGTCAAATGG GATCTCATTT	2160 20	ILCHAMBATO ILCHAMINA INTERPOLITA PARATEMENTA NATURATORIO GALGARATA	
	TRAGICCIOC CITICCOCACA CTCAGAACGG CANCOCCAAG GCCCGGAGTG TCCAGGGCTT	2220	CICCHARACE MITTERIORA IITORACIA ILLOGOMANA PALITRICII PARCALOGAMAN ANTOROGANAN PARCALOGAMAN ANTOROGANAN PARCALOGAMAN	
ď	стосостама отпалатетос смогоссама владосмом'я остинавляст сманальсес	2280	וראומסימאר אאוומסאראי מוזעראישט בעייואר וראימסימאר ביריסאימאין	1 5
3	CATTOCCACA GGCGGKGGGC CCAGCAGCAC CAGTGGAAGC TCAGCTGTCC TCCAGCTGCT	2340	TECTERATES CENTITIET ARTECTATES ANGACARITT ARTITIONS STANARIUS	77
	CTCGGCAGAC AGITCAGTGC ACAGITTATG CCCTAGCTGA AAAAGATCTC CCGGACGTAT	2400	Taracticac aggaecacit tiogradato citogragga arabigaras alogatrios	179
30	TTCAGGACAT COTCTTCCTC CTCCTCCTCA GGGCTCCTGC TACAGGCAGA GCTGGAACCC	2460 30	AAGAAATTAT TCAGAGAATT GAAAACGTTG TCCTAGATGC AAACTGCAGT AGAGATGTAA	132
	CONTRACTOR GREAT GREATHINGS GYCAGTICCT GTC	2503	AACAGAIGCT CTTGAAGCTT GTAGAACTCC GGTCAAGTAA CTGGGGCAGA GTCCATGCAA	138
	מרכסקרנורן מסקתמססקרו משפקרוסקי מו מיייים		CITCAACATA TAGAGAAGCA ACACCAGAAA ATGATCCTAA CTACTTTATG AATGAACCAA	144
35		35	CATTITATAC ATCIGATGGT GITCCTITCA CTGCAGCTGA TCCAGATTAC CAAGAGAAAT	150
	(2) INFORMATION FOR SEQ ID NO: 96:		ACCAAGAATT ACTTGAAAGA GAGGACTTTT TTCCAGATTA TGAAGAAAAT GGAACAGATT	156
ç	(1) SEQUENCE CHARACTERISTICS:	OV	TAICCOGGGC TGGTGATCCA TACTTGGATG ATATTGATGA TGAGATGGAC CCAGAGATAG	162
}			aagaagctta tgaaaagttt tgttttgaat cagagggtaa gggaaacag taaagttaaa	168
	(C) STRANDEANESS: GOUDLE (D) TOPOLOGY: linear		TITCAGCATA TCAGITITAT AAAGCAGITI AGGIATGGTG ATTTAGCAGA ACACAAGAGA	174
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 96:	45	GCAAGAAAT GTGTCACATC TATACCAAAT TRAGGATGTT GAGTTATGTT ACTAATGTAT	180
	CTGGAAAGCC GAGGGTAGCC GAGCGGGGCG GGCGCTCTGG AGCGGGGGGT GCTCGGGCTG	. 09	GCAACITIAA TITIGITIAA CACTATCIGC CAAAATAAAC TITATTCCCT AIAACITAAA	186
Š	COGITOCOCITO COCCAGAAGO ACCAAGAGAGO CGAGOCGAGG CCCGCCGCCC TCCTCCTCCA	120	atchchatat atalbadaa cittattaig tacacitaat ictacticitt tgcctgcaat	192
3	TCAGGCCCGA GTCAGGGGGG GCGGCTATAG CCCACCCGGG GCGCCTTCCC CCCGCGTCCT	180	AAAATCGAIT TIGAAAIAAA TGAAATGITG AAAATTITGC TAGTTGGITA GATGCTTATC	198
	ATCGCGAGCG CACGACAAGC GGCCCCTGGA GGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGA	240	CITIAAAITC TACTITICIT GAGGGGAAAA AGICTICGIC TGGAAATACA TATTACIGCA	204
55	ACCOTITICGA TCCGGCCCCA GGTGCTGGTC GGGGCCGGAR CCGGGGGCTG GGCCGCGGAG	300 55	AAAATGIAGC ATCCTTTTTT AGGIAGGAGT ATTATAGCTT YCATTTTAGT TKGACATTTA	210
*	GOGGCGGGCC TRAGGGCGGC GOTTTYCCGA ANGARCGGR GCCTGCTGAG CGGRCGCGGC	360	GIGTCCCAAT GAATTGAATT TCAAATATGA ATCATAATCT TGAAAATCTT TAGCACTAAA	216
99	ACCAGCGGC GCAACGCAAA GCCCGGGGT TYCTGCARCC ANCGCCGCTG CGCCARCCCA	420 60	GICTIGGGAA TATATCAACA ACTGATTAC ATATGCAGAT GCTATTTGNA TACCAAGGGC	222

AGGGTTCCCT TOGATCAGAC TCCTCTTTT TATCCATGGC AGGACTGGGC ACTTGGAGTC GODACTICCA AAATCCATGA GCTCTACACA GCTGCTTGTG GTCTCTATGT TTGCTGGCTA AGCCTCATCT GCCTTACTTT ACCAGTATTT GCTGGCCGTT GGTTAATGTC GTTTTGGACG AATTYTOCAC TCAGGATATT TCTGTTGATT GTCTTCATGT GTATAACATT ACTGATTGCC CACCAAGCCA TACTCCAGCA GOGAGGGCCT GITGGYTTTC AGCYTTACCG CCGACCTTTA AATCAGCATG CICGAAATAA CAACGCTATT CCIGIGGIGG GAGAAGGCCT TCATGCAGCC TCTTATTTAT TGGGAGACCA GGAAGAAAAT GAAAACAGTG CAAATCAACA AGTTAACAAT TESCIPAAGG GECTEGTECG AGCSTGGACT GTGACCGCCG GATACTTGCT GGATCTTCAT GAGCTGCTTC TGCTTCAGGT TGTCTTGCCA GCATTACTCG AACAGGGACA CACGAGGCAG AATTITICITIC CATACAATOT CATGCTCTAC AGTGATGCTC CAGTGAGTGA ACTGTCCCTC GIAAATAGIT CIGITAATAA CCCACIGITT TACATTIGGT ACATCIGIGI CIGCIAATAC ATGGAGCCAA AGACAATCAC TGATOCTTTG GCTTCTAGTA TAATTAAGAG TGTGCTGCCT AGTIAGCITT CICACITITC IGCITGITIG TICAGICIGA ATTAAAATTA GACTITIGAAA TCAAATCTTT TICCAGCTAA CTAAAAACTG TGTACAAAAG GATTGCTTGT AAATATGCAT TATTIGICAT ICAAGITITIC AICIGCITTA TAATIGATAC ACCITGAGGG ICACITIICI AACTGRCTAA CTTCATTACC TTAAAGCCTA GAACATTATT CTGCTTTATT TATATGGCTT AATACTTTTA CTATAATGTG GTACCACCTC AGCCCTAATA AATAATATTT TTACCTAATG TCTCACTITT ATTITIGTAGC AKGGGTTGCA TCGACTTTTT TACTAGAGAA TTITIACTAGA GACTICACIG GAAGATITAN ICCAANICHA GGAANIGTIC TITITTATIT TIATITITITC TTTTTAAATG TCATGGGGG GAAAAACCCA ACTTGGTGGA ACTCCCAGCT AAACAACCAA TCTCATGATC ATGAAGACTT TGATAGTTGC GGTGCTGTTG TOGTCATTOT GGCTCCCCTG AGGGACOCAG AGTGATCTTC 2801 2760 2520 2400 720 660 600 540 480 420 360 8 240 180 780 120 8 \$ 8 2 8 35 ઝ 25 8 5 5 S TCTTCCAAAA AAAAAAAAAAA AAAA GAAGCITICIG TICAATCCCA GCGGICCITA CCAGAAAAAG CCIGIGCATG AAAAAAAAAAGA CATGCTCATC GTATCTOTGT TGGCACTGAT ACCAGAAAACC ACAACATTGA CAGTTGGTGG GAAAGGCCAC GTGAAGATGC TGCGGCTGGA TATTATCAAC TCACTGGTAA CAACAGTATT COCAGAACCT ACTOAGGCAG CCAGCTGAGA AGAGTTGAGG GAAAGTGCTG CTGCTGGGTC CCGAGCTGGG CGAGAAGTAG GGGAGGGCAC GAGCCGCCGC GGTGGCGGTT GCTATCGCTT GTATTTGACT TOTOTTCTCA GCATTCAGAG AGCAGCGGTG TAAGATTCTG CTGTTCTCCC CGGATTIATC CATTITIACT GATGGTCGTG GTATTGATGG CAATTITIGTC CTTCCAAGTC AGGGGTGTTT GCACTTGTGA CAGCAGTATG CTGTCTTGCC GACGGGGCCC TTATTTACCG TGCAGACGCG ATGGATAACG TGCAGCCGAA AATAAAACAT CGCCCCTTCT GCTTCAGTGT GGTACCCAAA T TACCCTAAAA CCTTGGATTA AACAGAATOT GCATTGTACA TCTTTAAACA AAATGTATAT GCITCIGGIG TIGITICCITY ACTAGGIGIT ACTGCGGAAA TGCAAAACTT AGTCCATCGG agtittgtaa tittiatatta cittitagit tgatactaag tattaaacat attictgtat (2) INFORMATION FOR SEQ ID NO: 98: TAATTTATTA AATCTAGTTG TCACTTTAAA AAAAAAAAA AAAAAACTCG AGGGGGGCCC TOGATICTICT GACATTACTG CIGICIGAGA TITIGTATATO TOTABATACA AGTICCTIGA TGGACTICIC ICTITIGGAGA TITTITCCCAG IGATCICICA GCGITGITIT TAAGITAAAT CTCGTGAACT ACGAACGGAA ATCTGGCAAA CAAGGCTCAT CTCCACCACC TCCACAGTCA COCCAGTTTA AGCGCCTTTA TGAACATATT AAAAATGACA AGTACCTTGT GGGTCAACGA GTAATTGAAC AGGITTACGC AMAIGGCAIC CGGAACATTG ACCITCACIA TATTOTICGI CTGCATGCCA ANATCATTGC AGCTATANCA TIGNIGGGIC CICNGIGGIG GITGAMANCT ICCCAAGAAT AAAGTAGTIG TCTCAACAAC TIGACCITCC CCTITACAIG ICCTITIITIG AMACTIGACAG CTCCCGTGAT CTCTGTGCTG TTGCTTTTCCC TGTGTGTACC TTATGTCATA (1) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3 € (A) LENGTH: 504 base pairs TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear

> 1620 1560

1500 1440 1320

1200

1080 1020

960 900

1631

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CAGAAGGTTA AAGAGTGGTC

ACCATAAGGG CTGTGACGGT GATGGTGGCA TGGATGCCTC

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acroandria receiencer rendedere enertrande

480

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360 300 240 180 120

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO:

25

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INFORMATION FOR SEQ ID NO: 97:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1631 base pair
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear LENGTH: 1631 base pairs 20

ATAAAGCTTA AAAAAAAAAA AAAAAAAAAA AAAAACTCGA G

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WO 98/39448

320

PCT/US98/04493

322

WO 98/39448

PCT/US98/04493

GGCTAGGACA ATTITGGTGC TTTACCTATC TCTGCAAAGA CTGGAGAATT TGGCATACCA CAGTGGCCAC ICTCGAAAAA AITITAAGIAT CAGAAGAITA AAAAGAITIT AGGAITITGGA AGCITIGIAIT GICTITICCCC AATAATCAIT GITIGAICTC CAAATAGIAG CCITATAITA GCAATRGACA GATCATTICGT TCTCCATATIC TGATCATATIG TTACTACTIT GGAATCAGTA TTTGGGCAAA ITCAAGCAIT TAIGCAGIGG AIATAAAIGG AAATATAAAA ATAITIGCCA ACCIGICICA GRAACTITATIC ATTAICTICTICT GNATICCTICAA GGAAAGCACT TITIGCTITTTA CITTAGAAAGC GITICAGAIT IGCITITATAG ACTOCTIGCIG ICTICAGIAC CIGATAAAAC ITTAACCAGG GAAGCATTAA ACACAGTGCA GCAGCTTTTG CCCAGGCTTC TAAGTTCCTG CCGGCAGCAT ITATICAATIOT AAGAACTAGG ATGCTTCCTG CAGTGGCACT ACCTTCCCCT AGAGCTGGAQ **GTCAGTAAAA** CTGAAACCCC CKGATACTCA GACTITICALL IGATATCATT AAATOCAGGA CAGTOCCAAG AAGTOCTTGG AGTOTOGGCT CTGACAGOOC AAGAAGGGAA ATAACTTGTA TTAAGGAACA ACTATGAGCC AGGCCCTGAG CTGTCTCTA GATAATAAAA CAGATGGGGA GTGGAAGAGT CATTTGCTTC AAGTTATACA GCTAGGAAAT ACTICAAGICIA AATICITIGAAG GCAGICICICIC CIPATITICITIG GGACAGGICAC TITIGITACICAC ACACCATGGT CCACCTAAAA ACAGAAGGAT AAAAAGACTT CAGGTTTTCC CACTGTGTGC THANTIACAA CCACCAATCA TATCCAACAA AAGTACCCTA AAAGAAGGAC CTCCTCCCAG **OCAGGGTTTIG** TAAAGAGAAG TCACCAAGGG AGGCAGGTAA TGAATGTTTC CAGAATCAGT TAGGAATITIC TGGCTATCTT TCAAATGTTG AATTITCTGGA TGCTGAGAGG SEQUENCE DESCRIPTION: SEQ ID NO: 100: i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2847 base pairs
(B) TYPE: mucleic acid
(C) STRANDELNESS: A. '
(D) TOPAT CATGCTGCTT GOCCTTAAGC CCCAGCATGA TGAGGCTTCC GITTAGAGAGC TCAGAATTGG GTCTTGCCTG GGTGCAGGTG INFORMATION FOR SEQ ID NO: 100: ž 3 3 S 2 2 ೫ 25 39 33 45 <del>\$</del> S ŝ 22 8 240 300 360 420 480 540 8 99 22 780 840 8 980 1020 080 1140 1200 CACAGGCTGG CCTGCCGGTT CCACAGGGTA CAGTTAGGAC TTGAGTCTTT CTTTTTCTGT TTTGAGTTGG TGAGTGAGTG ATAGGGTAAC ATGGGCCTTC AGGATGACCC CTTGGAACTG TGCCGAGTTC CITIAANICIC AGCIGGANC CIGGACCIGG GAGOCCCCIG IGAGGGCCAG CICIGGAAAA CACTOGRAGE TETEGGGGTG GGAGGGAGAG GGGCTEGGGC TETETETGAA ATGAACACTG ACCTOGGAGT TGATGCCGGA GCTGTGGAAG AACTCTGCTC GAGGGCAGGG TGCCCTGGAA CICTICAGCA GITCAAGIAC ITGITCICAA AACAITITICI AAITGAITGG IAGGITITICA TANGCATTOT ITCTTTANGG CATGGAANGG GAAGAATGCT CAAGCAAGTC ATGTTTTGTTT TCAGTGGGAT GGCCCCCCT TCTCACTGCT GGGGCTTCC CCTTCATGTG GCACCTTTGT GCAGGGGCCA CCAGGCAGAC TCTTCCCACC TTCTCCCACT GAAGCACCAA GGGGCTTGGA ACCGRANTITY GGCTAATCAG AGGCATTITY TITIGECCIAG TATCTITICAC ACTIGECCAA COGNOTIATY ITITIAAAG INCIOTIGOT IGTANTAACA CGAAACTAGA GAGAAATAGI TICTGAAGCC AGTITATIGI GAAGAICCCC AAGGGAAGGI ICGGIAGAGA AAAAIAGIAA GCTGGTTTAG AAACTGACGA GGGCAAACAG CCAGGACGCA TTGGAGAGGA ATTTGCCAAA GATCTACCCT GAGATAACGC CTGTCCAGTG TCTTCACCAC GTGAATAACC AGGGCTCCAA TTTAAGAATC CATGTGACTT TAGAATGGAA CTGCCGGCCC TGGCAACTGT CACGTGTGCT AGAAGGTTCG THAGATTGAG CTGGGCAGCT GCAATCAGCT TCTTTATATG CAAATTAGGC ACGACCCATC ATGCCTCTGG AATGCATGTG ATACTCATCT CCATTITIGIT TCCTTGAITIG CATTITITIGIT CTITITAGCAG ATCTGTCCCT GTGGGTGGTG TCTAAGAAGT CGGACACCTT GGTTTTTGTG

260 320 380

TGACCATCCC AATITIATGAA TCTTCTTCAA AATGACATTT CACAGITATA GITAGGGCTC AGABATGGCA TTGAGGTAGC CTTATTTCTC CCCTTTAGCA GATGCTTTAA GTACACATTG CTGACTTGAG CCCACCCCA GGAGTTAGGA GAACATTTCC TTTTTCATGC CATCTTCCAT

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1260 1320 1380 1416

IGTOGITICCT GÓTTGGTGGC TAATGAAGTG AGGGAGGGA GGGATGTCAC CCCAAAAGTA

AGTOTITITIC TOCTITIGAAA AAAAAAATTC CACAAGCTTT TAAAAGGTGCA

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S

TIGOCITIGO CCAGGCCAGA CACITCACAT COTTIACATO GITCIGIGIA THAIGTGTAT AAAGCGAAGC TGTTTCTGTG AAACTGTATA TTTTGTAAAT

OCCCTCCCA ATTITITABAGT

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AAATATATIG CTACTIGAAA AAAAAAAAA AAAAAA

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AAATAAGGTG TITCTTGGCC TICAAAGATA TAGAACTTTG CAGCAGTAGT AAAAGTGAAG

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980 140 200

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GGCACGAGGG AGGGAGCCT CTCCCTTGGG TCACTCTTGT GTGCCCTTTA

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SEQUENCE DESCRIPTION: SEQ ID NO: 99:

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.) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1416 base pairs
(B) TYPE: unclaic acid
(C) STRANDERNESS: double
(D) TOPOLOGY: linear

(2) INFORMATION FOR SEQ ID NO: 99:

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WO 98/39448

PCT/US98/04493

324

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 794 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

8

S 2 INFORMATION FOR SEQ ID NO: 102:

> 1380 1394

1320 1260 1200 1140 1080 1020 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 8

TGCTGATCAC ATCAGATTTT TATOTTTAAA AAAATCTCAT TATOGATTGA GTCCAGCCCA CATATAATGG CTGTGCAATA CATGCTTCTC AATAAGAAAA TTAACTGCAT GTTTACTGTG CICCAGITAG ACCTAAGOGC ACAAATOCAG AATTCATGAC CITGIAGITG TGGCAGGGIC TAAGGCACTG AGTCAAAGTG ACAGCCCTGA AGGAAATTGC ACTCCAGCCC TCCTCCAGGA GTTGGGGCTT AGGTACTIGC TTACAGGAAG AGCAATTCCC TAGCAAAGGT CATTAGCTCC ATTATICTIOC GTACTACTIC AAAGATTICT GICAGCCCTA ATTACAAGIG ICACCATATA CATOTICCACA CACACACACA CAATATTTGA GAGCTAAGGA AAACTCAAAG CAGCCCCTTC TCAAAGICCA GIGAAICIGG CICICITACI GATICCIGGI TITAGIGIGI GIGICGGGG TAAATTATIT TOTAAGAGAG ATTTACTOCT ATCCCAGGAT GITCGGACIT GOTGCCCCTG GTTCTTGATT CTGGAGGCCT GCCTGGTAAG ATAAGATAGT ATAATTTTGGA ACTGAGAACA CCAGCAGTGC TTATRAAGCC CCCTACCCTG TCCCATTCCA GAAACCATAA GACTCAGGCA GCTCTAAGAG AAAAAGAAGG CCCATATGGG AGACTTCAGT CTCATTATTA TTGCCTTTAT TOTCTAATAA GATGGGAAAC TTGGATGCCC AGCCATTTTG GTGACCTGAG AGTCTAACTA AGTGTGTACC TATATATAAA GGACAAGTGT GATATGTGTG TATATGTATA TACATACATA NAMANACTO GAGGGGGGCC CGTACCC TOCATTTOGA AATCAATAAA CTATTACTOG AAATGCCAAA AAAAAAAAA AAAAAAAAA TGAATCTTTC AAGGGAATTA CACGTTTGGG TTAATGTTTC AGTATATCAT TTTCATACTG AGICACATIC CCICCITAGO AATCITCCCC TICCACCCII TACATIAAAC AAGGGAACAC TACCAGAAAC AGCAGAACGA GGGCCAGAGC AGAAAAAATGA AAATAAGTOG AGACACTTNT ACCAMATETT CATATTTIMG TAAAETTAGE COCCACTGTA CTCTGTGAGG ATGTGCCAMT TCTCTCCCCA AGTAGAAAAT ATTCTCTTGC CATTCCTGAA ATTCCACATT ACTITICAGG 2160 2100 2820 2700 2640 2580 2520 2460 2400 2340 2280 2220 2040 1980 1920 1860 1740 1680 2847 2760 1800 1620 1560 8 3 6 35 မ 25 20 5 5 S GCATTGCTGC ATCCTGGGAA GGGGGTATAT GGTCTCACAA GTTGTTGTCA TTGTTTTTTT TOTOTOCAMA ACTOCAGAMO CTCACTOCCT ATAMGAGGAA ATAMGAGAGA AAGTOGAAGA AAAAAAAAAA ACCC GCATGCTTTC TTAATAAAAA AAAAAAAAAA ATGTTTANAG TTTTATCTTA AAAAAAAAAA GICCCIOCIC CICAIGITIC IGGITIGGIG AGICCITIGI GCCACCACCC ATAAIGCTIT GAGGGACAAA AGGAGTAATT ATTTGGTATA GATCCACCCA TCCCAACCTT TCTCTCCTCA TITICTIGITIG GICTAGCATT GCIGGACACA AAGIGTAGIC ATTAITGITIG TAITIGGGIGA COTGACTOCC TITIOGGOODE AGITTICCOCT COCCTICATG AAATGAAAAG AATACTACTT NTCNCTUAGA TICCAGCICC CTICCACCAA GCCCAGICTT GCTACGIGGC ACAGGGCAAA TICIOGCOTO CAGCAGGOTA GCIGAAGITIT GOGICIOGGA CIOGAGATIG ATCATACATO CAAATOCCCC TIOTICATCI OTGICTICIO CAAACTAGIC TCATGAAGAA GOOTOTOTICC TATACAAAAC TTCCCCATCAG TTCTCCCCAA TATTCCCCAT TTGTAAATGA AMAGAMATOG GGOTGCGAGT GGCTTGAATC TCCCATGATG TTGGAGGGCA CTTAGTGGGG GAACAATTIT ACTICIGICC TIATITICACI IGCIGAAAAG CIGIOGGACA AAAIGTAIGG TCACTTCTCT AATGCCTTTG CCCAGTGTCC CTATTTTAGG CATCTTTTCC TICCTTATTC CTTCCAGTCA TTCAAGTATG ACATAATATT TCCCATTGGG GAAAGGAGAA TTTCTCTTAG AGGGTGGCAA AAAGGGGCAA GATAGGGCAG TTAACTAAAG AGCACTTTAT TTCTTTGAAG CCTTTCTAAG CTITITITIGG AAGGGGGTTA TATATGAGAG TICATTGAAG AAGTCCAGTG AGGCTGAAGT ATAATTICCA AGAGGITIOG COTTCCGCIC TCCTGCTTTT TICTTICATC CACCCCTTTC AATAGACAAG GCCACTTTCT TIGTGATTTC TGCTTTTCAT GCATATTATT TTATTTACCC GTTGCAAAGT ACTCAGTGTA TATTTAATGT TGATTGTTGA ATTTTAGTTA CGAGAGGGAA GAGATTOOTG GAGGAGAGTA AATAATCTAG AGGCAAGAGT TCAGTGAGGG CCAAGGGGGA CCCCCAGAAA AAGGTATGGA GCTAACTCAT CTCTTTTACA AGGGGTGGCC ATGACTTACT TITICTAAACC CITITICCTOT TCAGATCCAT ACAGGATTIG CAAGGGTAGG

GCCATTAGGC

GGTGTTCTGC TCTCTACTCA ACTTVATTTG AAAATGTCTG CAGCTTCACT CCTGTAGAAA

1500 1440

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:

101:

WO 98/39448

323

GACAGCAGAT TAATACITAA TGAGGGTTAA ACCTGACCAG TCTTTCTACA GTGACAGGCC

GGATACATTG GTGCAAAAAA AGCCACGGGS CCCATACTGG GCTTGATATG

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ACACTOCATO AATOGOGAGA ACCAATGAAT CCATTOTCCT CTOCCTATTT TCCTOTOCAC

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INFORMATION FOR SEQ ID NO: 101:

SEQUENCE CHARACTERISTICS: TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

WO 98/39448

326

325

linear
TOPOLOGY:
9

(xi) SEQUENCE DESCRIPTION: SEQ 1D NO: 102:

•	GENRICANGGE GEAGTANAGE GACTICANGES AGECRATICE CEGATIATIC TATTICECET	9
	CCCTCTCTCC CGCCCGTAT CTCTTTTCAC CCTTCTCCCA CCCTCGCTCG CGTACCATGG	120
9	CGANGCOTCG GCGCCACTC AGTCCCATTC CATCTCCTCG TOSTCCTTCG GAGCCGAGCC	180
2	STCCCCCCC GGCGGCGGG GGAGCCCAGG AGCTTGCCCC GCCCTGGGGA CGAAGAGCTG	240
	CAGCTCCTCC TOTGCGGTGC ACGATCTGAT TTTCTGGAGA GATGTGAAGA AGACTGGGTT	300
15	TOTICITITICA CACCCIGATC ATOCITICATI CCCTGGCAGC TITICAGIBTC ATCARTGIGG	360
	GITTCITANG TCATCCTGGC TCITCTCTCT OTCACCATCA RCITCAGGAT CTACAAGTCC	420
5	GICATCCAAG CTGTWCAGAA RTCAGAARAA GGCCATCCAW TCCAAAGCCT ACCTGGACGT	480
3	AGACATTACT CTGTCCTCAG AAGCTTTCCA TAATTACATG AATGCTGGCA TGGTGCACAT	540
	CAACAGGGCC CTGAAACTCA TTATTCGTCT CTTTCTGGTA GAAGAICTGG TTGACTCCTT	009
22	GAAGCTOGCT GTCTTCATGT GGCTGATGAC CTATGTTGGT GCTGTTTTTA ACGGAATCAC	099
	CCTICIAAIT CITOCIGAAC TOCICATITI CAGIGICCCG ATTORCIAIG AGAAGIACAA	720
S	GACCUAGNIT GAITCACTNIG ITGGCATOSC CCGAGAITCAG ACCAGGTCAA TIGTIGAAAA	780
3	GATCCCAAGC AAAA	794

(2) INFORMATION FOR SEQ ID NO: 103:

35

(A) LENGTH: 1544 base pairs
(B) TYPE: nucleic acid
(C) STRANDELMESS: double
(D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: 6

TITICCTIGCT AGICIGAACC AAAGAGITGT ITGGGCAITT GCIGIGITIGG CCATITICIGG CTGGGTGGAA AGAAGTATAC CTTTCCCTGG GGCCCTAGGA TAGCAAAGTG AGCCATAGTG AGCAAGAGGG TCTTCTTCCT CCTTCCCCCA GCCAGCCAGC TGTCCTGGGG CCAGGCTTTC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103: S 45

240 300 360 420 480 AGCTATGACC ATCTGTGCGG TCAGGGTACA CTCAGCTCTC CTCCCCAACT CCAGCAGCCT CTCCCACAGG GAGGACGGAG CTGGCCTCAG GAGTGGGACA CCCAGACTTG GCAGGGCTT TTTGAGCCTT AGCCATCTCC TGTCAGGTAG GAATGAACTT GCCAGCCTTC AGGYTCGTTC TTAAGAAGTG TCCCTTTGGC GCCCCCTGGA GGCAGAGCAC TGAGCTGGAC CCTGGGGTAGA GOCCAGGOTO COCTOCATOC TOGOCOCAG COCAGOTOTO CACTOGOOTO GATCACOTTO 8 55

540 900 99 960 070 906 080 140 200 440 1500 260 320 CAAGAGGCCT GTGTGGGGGC CCCAGGAATC CTTAGCTGAA GCGGGGAGAC TCACTCTCA TCTCAGGAAA TTCTAGCCCT TCCCCTCAGG GAGCCACGGT TCAGGGTGAG GCCCAACACC ACTCAGGCCC ACTICATTICCT ACACATIGCC AAGAITICAC ACAIGIGACC AGGGCCACC AAAGICCCTG IGACCITIGI GACTAGGATC CTAATITICITC TATTITICITC TEGGIGCCTG GENCTIGITGTC ACCTGGGGCA GIGTOGATAA TOTITIAGITIC TOTCACACTO TITTITIOOGO GIGGCACCTIC GITCTCCGAT TAATGAGTGA TOCTATGGCT TOCTCGTGTC TTATGATCCA ATCCTTTTCT ACATCAGCCC OCCIGOCCIO GIGICAGOCC CAGGACTOTA GIOCTOGGAG CAGTAAAGCT CAGCICTIGIG INCITITICIT ITALCOCIAG ICITIALCIGO CCIGOITAIT ICCTIGCOGG GAGGAGAGG TITIGETABLE TECTECCAGE CEAACETAIT ACEAECECAC CICECTEGGA CETACTGCTC GOGAGOCAGO AGACAGGAA CCACCAGOAG TGGCTTCCTG GCCCTGTGCT GGGGGTQQGG GGAAGCTGGG GGCACATGTG GCCCTTGCCT TCTGAGCAGC TCCCAGTGCC AGGCTTTGA GACTITICCCA CATGATAAAA GAAAAGGAG GTACAGAAGT TCCAATTICCC TITTITATTIT GCTGGTTGGT ATCTGTAAAT GTTTAATAAA TATCTGAGCA TGTATCTATC AACGCCAAGA ATTICADAGT CICCITICADE AATAIGAGGE TITITAGGAIG TITATATICE TICATECETIC ITGITITCCCA GGITITICCAG GGAAAAAAG ICIGGAAITA IAGAIACAGC ITAITAITAA TOCCTTAGGG CCCTGGGTGG GCAAGTCTGG GCCCTGGGGT AGGGAGGGAG ACACTIGGG AITITICIAAT TICAGACAAA CACACACTCA GCGCGCACTC ATTIGITICIT GCATAAAAA AAAAAAAA AACNCNNGGG GGGG S 2 2 ន 23 9 35

(2) INFORMATION FOR SEQ ID NO: 104:

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(A) LENGTH: 871 base pairs TYPE: nucleic acid STRANDEDNESS: double (i) SEQUENCE CHARACTERISTICS: æ <u>0</u> 0

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TOPOLOGY: linear

ACCCACGCGT CCGNCTTGTC CACCCGGGG CGTGGGAGTG AGGTACCAGA TTCAGCCCAT TIGGCCCCGA COCTICIOTY CICGGAATCC GGGTGCTGCG GATTGAGGTC CCGGTTCCTA AGGTGGGTCG CTGTCCACCC GGGGGCGTGG GAGTGAGGTA CCAGATTCAG CCCATTTGGC CCCGACGCCT CTGTTCTCGG AATCCGGGTG CTGCGGATTG AGGTCCCGGT TCCTAACGGA CTGCAAGATG GAGGAAGGCG GGAACCTAGG AGGCCTGATT AAGATGGTCC ATCTACTGGT (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:

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9 120 180 240

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WO 98/39448

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PCT/US98/04493

50 25 6 35 30 23 20 7 5 AATAATOTCA CTOTOGACCC AGTOTCACTC CTCCACCCCA CACACTGAAG CAGTAGCTTC CCACTITCCCC COOOCAGCTC CAGGGATGTG GCCTCATTGC TGTCTGCCAC TCCAGAGCTG TOOGTCAGTG CCCTCTCCAC CTACACCTGT GACCAAGGCT TATGTGGTCA GGACTGAGCA CACCATCTTC TTCATGCTGT TCTCCAACTT CTGGTATCAC TCTTATACCA AGGGCAAGCG CATGACAGCC ATTCAGCTGA TOCAGTTTGT CCTGGTCTCA CTGCACATCT CCCAGTACTA CAGCCCTGAG GCACTTAGGA TOGITICGGGT GGCCTGGCTC TTCCTCTTCT CCAAGTTCAT GGGGGGCCCG TACCCAAATC GCCGGTATGA TCGTAAACAA TC GGGGCTAAAA GGGCTGTACA GTTATTTCCC CCTCCCTGCC TTAAAACTTG GGAGAGGAGC CTTTATISTICC AGCTOTAACT ACCAGTACCC AGTCATTATT CACCTCATCT GGATGTATGG GGGAGGAATG GGCTCTTTCC ATGCCATGAT AAACTCTTCC GTGCATGTCA TAATGTACCT CITEGITICIE TEACITIGGE CIEGEATEAT GGETAATEGG AAGECETTEE AGCIECGIGG GATOGAGGCT GTTGTGAACT TGTACCAAGA GGTGATGAAG CACGCAGATC CCCGGATCCA AACTTGTGTC TTAATTAAAA GTGACAGAGG AAACCANAAA AAAAAAAAAA AAAAACTCGA TGGGCCAAAG GTCAGGGTGG GCGGGGGCCT GGGAATACAG CCTGTGGAGG CTGCTTACTC ACTICAGGGOT GGCCCCACAA AGGGTCTCGT GGCCTTTTTTC CTCACACAGA AGAGGTCAGC GOOGLACTIOC CCTCCCCTCC CCACAGCTGC TCTACAGGGA CCACGGCTTT GGTTCCTCAC AGAAGCATOO CCTAGATAGG COCCCACCTA AGTOCCTCAG GACTOCACCT TAGGGCAGTO GCTGCCCCGT GCACTTCAGC AAAATGGAGC TCCAGGTATT GCCAAGGTCA AGGCCAACTG GTACTACOGA TTATCTOCCT TTOOCCCTOT GOCACAACCC TACCTTTOOT GGAAAAAAGCA ACATOTICTIC CATCACTCTO TOCTTCCCTG GAGCTGGTGG TGGGGGGTAA AGATTGCCCC TOAGCTGATG GACACAGTGA TCTTTATTCT CCGAAAGAAA GACGGGCAGG TGACCTTCCT CCTGATGTCG GGCTGGCTGA GCACCTATAC CTGGCGCTGT GACCCTGTGG CTICATGATI GICTACAACI ICICACIGGI GGCACICICC CICIACATIG ICIAIGAGII GOOCTACCCT CTGATGOOGT CCCCCTTGCT AATGACCTCC ATTCTCCTGA CCTACGTGTA GOCTICTRICGA GAGGAGAGAG GAGAGCOCTG GAGAGGAGAG OCTOGAGAGT CCTTAGCCAG GTCAGACAGG TOGAGCCGCC GGGGCAGGAG TCTCAAAGAG CCAGGCTCCA GGAGAGGAAG ACTATTCCAA 1500 1440 1380 1320 1140 1260 1200 1080 1542 1020 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 6

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 106:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2327 base pairs (B) TYPE: nucleic acid

INFORMATION FOR SEQ ID NO: 107:

LENGTH: 1542 base pairs TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 106:

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AATTIMTAAT NIGAAAAATT TITTIGGGGT TITTIGGGGCC ATGG

300 360 404

TCTAAAATTT ATTTTTTTAA AAAGAGAAAC TGCCCCATTA TTTTGGTGGG GTTGGTTTTT

TITITAATAA GAATGACGCC CCACTITIGGG GACTAAAATT GIGCTATIGC CGAGAAGCAG

8

AGATACCAGA ATOGOTTACA CATTIVAACCT GOCAAACATT GAAGAACTCT TAATGTTTTIC

240

120 180

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AACTOCTTTG AGTTACTCAG AACTTCAACC TCCCAATGCA CTGAAGGCAT TCCTTGTCAA

GOCACGAGTT ATAGCATGGC ATTICATACTT TTGTTTTATT GCCTCATGAC TTTTTTGAGT
TTAGAACAAA ACAGTGCAAC CGTAGAGCCT TCTTCCCATG AAATTTTGCA TCTGCTCCAA

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LENGTH: 404 base pairs
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 105:

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 105:

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AATGCTTCTT CAGAAAAAAA AAAAAAAAA A

360 420 480 540 600 660 660 720 780 780

2

TOGECACTOT CAACGECCGE TOGETGGAAC CECGEACCAC AGETGCEATG TOGGCCTGC
AAACCCTGGG AGAAGGAGCG AGGCCTGGGT GGGGAGGTAC CAGGCAGCCA ACAGGTTCCC
GATCCTTAAC GCCAGNTGGG AGAGAAGGAC CCCAAGTACA GTGCTCTCCG CCAGAATTTC
TTCCGCTACC ATGGGCTGTC CTCTCTTTGC AATCTGGGCT GCGTCCTGAG CAATGGGCTC

TOTOTOCOCTO GCCTTOCCCT GGAMATAAGG AGCCTCTAGC ATGGGCCCTG CATGCTAATA

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TECACATETE CATOGGETOT GECTTEATEA ACCTETIGAT CITIGGETTEA CAGEATGETT

CTIGICAGOT GCCIGGOGCA IGCAAAIGIG GOIGACCIIC GICICAGGCI IICCIGCIITI ICCGAAGCCI ICCCCGACAI ACCIICGGAC IAGIGCAGAG CAAACICIIC CCCIICIACI

GOGETEAGET CACATTETOG GAGGECAGEE AGEITTAEET GETOTTEETG AGEETTAEGE

PCT/US98/04493

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2280

	(C) STRANDELNESS: double (D) TOPOLOGY: linear			TITIAACAATC GACAAATGAA ATTCTTACAG CCTGAAGGCA GACGTGTGCC CAGATGTGAA
v	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 107:		v	AGAGACCTIC AGTATCAGCC CTAACTCTTC TCTCCCAGGA AGGACTTGCT GGGCTCTGTG
`	GOTAGCTCAN TOCAGTGAAA TAGTCTTACT GGAAACAAAG CCCTTTATCA AGAATAATTA	09	ר	GCCAGCTGTC CAGCCCAGCC CTGTGTGTGA ATCGTTTGTG ACGTGTGCAA ATGGCAAAGG
	ACTICITICCCT TITICITITIG GAGAGGIGCT TIGITICIGA TEGGACCAIT TEACTGCAGE	120		AGGGGTTTTT ACATETECTA AAGGACCTGA TGCCAACACA AGTAGGATTG ACTTAAACTC
0	AAGCAACACA GTATTCTRAG CAGAAGATCG GGACTTGAGG CCATGTTGCG GAGGGCCAGT	180	01	TTAAGGGCAG CATAITIGCTG TACACATITIA CAGAATGGTT GCTGAGTGTC TGTGTCTGAT
	RACATTATET GGACTETGGA GTGTGAGGAA TATGGACTEC ACTETTCACT ATAITCACAR	240		TITITICATICC TOGICATICAC CTICAAGGAAA ITTAITAGAC GTATAATGTA TOTICTIGITGT
Ž	CGATTCAGAC TTGAGCAACA ATACCAGTTT TAGCCCTGAT GAGGAAAGGA GAACTAAAGT.	300	<u>.</u>	TITINACITG ATCATGATCA GCTCTGAGGT GCAACITCTT CACATACTGT ACATACCTGT
3	ACAAGATOTT GTACCTCAGG CGTTOTTAGA TCAGTATTTA TCTATGACTG ACCCTTCTCG	360	3	GACCACTCTT GGGAGTGCTG CAGTCTTTAA TCATGCTGTT TAAACTGTTG TGGCACAAGT
	TOCACAGAGG GITTGACACTG AAATTGCTAA GCACTGTGCA TATAGCCTCC CTGGTGTGGC	420		TCTCTTGTCC AAATAAAATT TATTAATAAG ATCTATAGAG AGAGATATAT ACACTTTTGA
20	CTTGACACTC GGAAGACAGA ATTGGCACTG CCTGAGAGAG ACGTATGRGA CTYTGGCCTC	480	20	TIGITITICIA GATGICIACO AATMAATGCA ATTIGIGACO TOTAAAAAA AAAWAAAAA
	AGACATGCAG TGGAAAGTTC GACGGAACTC TAGCATTCTC CATCCACGRG CTTGCAGTTA	240		ACTICAAGGG GGCCCGGTAC CCAAATCGCC GATATGATCT AANDATC
35	TICTTGGAGA TCAATTGACA GCTGCAGATC TGGTTCCAAT TTTTAATGGA TTTTTAAAAG	009	2,	
}	ACCTCGATGA AGTCAGGATA GGTGTTCTTA AACACTTGCA TGATTTTCTG AAGCTTCTTC	099	}	(2) THEOREMAN TOTAL TOTA
	atattgacaa aagaagagaa tatctttanc aacttcagga gtttttggtg acaganaata	720		(1) CENTRACT CHILDREN CONTRACTOR
30	GTAGAANTTG GCGGTTTCGA GCTGAACHGG CTGAACAGCT GATTTTACTT CTAGAGTTAT	780	30	(1) Selveral Christians (1) Selveral Strucks (1) LENGTH: 102 base paixs
	ATAGTCCCAG AGATOTTTAT GACTATTTAC GTCCCATTGC TCTGAATCTG TGTGCAGACA	840		(p) TYPS: MUCLERC ACAC (C) STRANDENESS: double (D) TOPOTICES: 1 tnear
35	AAGITICITC TOTICCITGG ATTICCTACA AGITGGTCAG CGAGATGGTG AAGAAGCTGC	006	35	(x1) SEDITON'S DESCRIPTIONS (DATE OF THE OFFICE OF THE OFF
}	ACGOGGCAAC ACCACCAACG TTCGGAGTGG ACCTCATCAA TGAGCTTGTG GAGAACTTTG	096		1007 ON THE RESTAURT SAME SERVICES STREET
	GCAGATGTCC CAAGTGGTCT GGTCGGCAAG CCTTTGTCTT TGTCTGCCAG ACTGTCATTG	1020		STATEMENT STATEM
40	AGGATGACTG CCTTCCCATG GACCAGTTTG CTGTGCATCT CATGCCCCAT CTGCTAACCT	1080	40	ANALYMENT MOONTHEM CHICKIEN CHAMBELLING ICLANGISING COLORES
	TAGGAAATGA CAGGGTTCCT AACGTGCGAG TGCTGCTTGC AAAGACATTA AGACAAACTC	1140		MACAMANA INSCRIMINSI UCHACINIAN IIICICINAN OSITCINAN GAIRCICITC
٧٧	TACTAGAAAA AGACTAITITC ITIGGCCTCTG CCAGCTGCCA CCAGGAGGCT GTGGAGCAGA	1200	45	CICATERIOS SECUCIONES CONTENIOS SE CONTENIOS CONTENIOS DE CONTENIOS DE CONTENIOS DE CONTENIOS CO
}	CCATCATGGC TCTTCAGATG GACCOTGACA GCGATGTCAA GTATTTTGCA AGCATCCACC	1260	!	THE STATE OF THE PROPERTY OF THE STATE OF TH
	CTGCCAGTAC CAAAATCTCC GAAGATGCCA TGAGCACAGC GTCCTCAACC TACTAGAAGG	1320		CANAGEMENT INCOMMENTAL MODERN PRINCIPAL MANAGEMENTS INCOMMENTED TO CANAGEMENTS.
20	CTTGAATCTC GGTGTCTTTC CTGCTTCCAT GAGAGCGGAG GTTCAGTGGG CATTCGCCAC	1380	20	GENERAL TOTALITATI TOTALITA CUITOCUCA TOTALUCATU ASACTUADA
	GCATOTICACC TOGGATACCT TTCCGGGGGGG GAGAGACCTT CCTCTCCTGC GGACTTCATT	1440		CARANCETICI COSTITITOS AGAGATIGAS GAGIOCACA TOCACTICOS TOTOCAAGGG GACAACTAG GOTTOCOTAS TRANSPORTAN CONTAGORAC CAMMINATAS ASTRA
٧	GCAGGTGCAA GTTGCCTACA CCCAATACCA GGGATTTCAA GAGTCAAGAG AAAGTACAGT	1500	55	CECTABLE WYCESARS: ACAMARICA CONTRACTOR CONTRACTOR
)	AACACIATT AICTTATCTT GACTTTAAKG KKMAMKMAM KCTCAGASRA TTATAMITISM	1560	}	SUCTIVITIES DESCRIPTION PROPERTY PROPERTY PROPERTY PROPERTY PROPERTY.
	CHIPHRARGSH WIMAANSCITK SHGCTCYNCC KSRSTGRMKG MIRCICCINGA AYITHGIRGAK	1620	4	CONTRACTOR CONSTITUTES INTOTOTICAL TITUTATES TO TOTAL CONTRACTOR C
8	CHYYYKSGCT KOMGGAAKKS GGCASGAGCC AGAGACCTGC ATTGCTTTTCT CCTGCTTTTTA	1680	8	MARKATATA CHARACTER CONTRACTOR CO

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1062 1020 1080 1020 960 900 840 960 900 840 780 720 660 600 540 480 420 360 90 180 120 780 8 3 S 50 8 35 30 23 8 5 5 S 2 ATAMACMSCT TCACAATTC TCCATGACTC TTATATACTG CCTCATCTTG ATTTATAAGC AAAACCTGGA AAACCTACAA ACAGAATCAG AGGCCATOGA TACTGACAAC TGATTTOTCT GTTTTTTTTC TCTGTCTTTT CCTACATETT CCTAAGETTT TTAGEAGGTA TATGTTGAAC ACTICTGTTT CATGGTTGAG GTTAAAAATC TATACTOGAC AGTTACAAGA AATTACCOGA GAAAAGCTTG TGAGCTCACC AATTWOGSAC AAABTBGHGG GGGDTCCAAA CHTWVTCGHG KAAMTTCTCT WAARMATYTK AAAATCAATT TIGIATAGIT TATTICAATC TAAATAAAAT GIGAATTIIG TIWWATTAAA **AATAAGTOTT** TIGAANGGAA AAGCCIGCIG TIGTICCACA TCICGTIGCT GITTACATIC CITTIGIGGAG AAACAAGGAT TICAGIGIAG ATTITIGICIT TCITGAACIT AAAGAAACAA AIGACAAAGI TTTATAAAAT GITGIAGIGA AGCCCACAAT IGACCIIKGA CIAATAGGAG IITTAAGIAT GAGAACGITG AAAGIGCCAI GITTICCITIT GOGIGATICIC IGITGAIGGG ACTEIGGAAT CIGITTAAAT GOCCCCIGIT TGAACICICA AGCTITGAAG ACCTACCIGI TCTTCCAGAA AATCACTAAA TATCTTTGCC TATAGGACTC CATTGAATAC ATTAGCCATT GATAATCTAC TAMARCACGA CCTGGATTTA ATOGTGGACA CATATATTAA ACTCTATACR AKTAMOTCAG GGAACACTTT GACAGACCAA AGGTCAAGTA ACTTGGCTTT GCTTAACATA AATTTTGATA TGATGAAGGT TGAGAATGAG CGGTATGAAA ATGGACGAAA GCGTCTTAAA GCATATTTGA CTGACATCAA GITTITITICCT AAIGIGIAIG CATIGCTGAA GGICCIGIGI ATICTICCIG GAAGTGACTT ACCCAATCCT GACACGCTGT CAGCTGAGCT TCATTGTTGG AGAATCAAAT TOGOTOTTAT GAATTCTTTG AAGAATATAT TTTGAAGAGG TOTGGGAGGA AGGAATACAT TOTTTCCAGT TTAAKTCATT TTAGACATAG CATTTATTAT CACTGTGGAT CTCTACTTGT COMMITYYMK RKTYGAYMYW YOCGWMCGAG AAAAAGCCGT AAGGTGTATO TAGACCACTT MICTYYCTAC AKAYRAYTCM SWAWMTOTOG AAARYWSSTA MIDWSWOCWKK TAMMRRIWCG GEAAACACAG GOGGAAAGAT ATAGAGCTTC COTOCACCAT CTATGAAGCC CTCCACCTGC CCTCAGTCAT GGGACAACTC AAATTCAATA CGTCGGAGGA ACACCATGCT GACATGTATA INFORMATION FOR SEQ ID NO: 110: (i) SEQUENCE CHARACTERISTICS: GIGGITTATC TAGAAAAATA TGGAAAATAT TGCTGTTATT (B) TYPE: nucleic acid
(C) STRANDEDNESS: double (A) LENGTH: 1751 base pairs TITIGGTGAAG 2539 2520 2460 2400 2340 2280 2160 2100 1200 2220 2040 1980 1920 1860 1800 1740 1440 1380 1320 1680 1620 1560 1500 1260

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TACAATKAAT YWTRRYTTSM KRMAGMYAAT CCGAAAYTGT GGMAAMYAAA CTTGATATTC

CCCTGGGAAA TTCCGCAGAG CTCACCAGGG TAACTTGGAA TCTCAGCTAA

ARMTTTTGAY ATRMARYACT RMMTKSAYTY AAYGRWGTGA CWSGAWAATA TTRAASTYTA

8

AACTTAAAGA TATATTCTCA GAACAGCACC TCAAAGCTCT TAAATGCTTA TCTCTGGTAC CCTCTGAGAG TTACTATAAA GAAACCCTAA GTGTCCCCAAC AGTGGAGCAC ATTATTCAGG 50

ACAAGAGCCT TIGGGAAAAA CYYCMAGGGG CAAACCICIG ATGICTICIT TOCKOMSRI

GCAGIGICAG ATTITIGATIT CATIGITACT ATTGITGITC TIAAAAAIGI CCIATCITIT TAGTGACACA AATATTAGAT GGAATAACTA TATAGCTGGC CGAGCATTTG TACTCTGAGT TOCTTTTOAA ATTITAGTOG AACTCCTGCA AOCACTTOTT TTATOTTTAG ATOGTATAAA TAAAGAAAGG GGTAAAGAAC TGAAGGAAAT CTGCCATTCT CAGTGGACAG GCAGGCATGA 35

30

AGCCTICCIG CCTIATGAAG CCGATGCAGA AATTITGGCT GTGAAATITC ACACTATGAT GCACCTACCT GTGTTGGTGA GGTTTGTTGA TGAATCTCAT AACCTAAGAG AGGAATTTAT GAGAGACTCA CACTICITIT CCATIATCAC TGACGATGIA OTGGACATAG CAGGGGAAGA

8

TOGATOACOA CAACTOCTITI TAGAACTIGA CAACGTAATI ICIGITICTITI TICAGAACAG

CCYGROOTOS REGIAWYTSK TOCAYKAGOG AACAATTGAG GAAGITTOIT CITTITITCCA

TOGATITICT TCCAAAATGA AAOTTOTTOC TICTAGACIT TYAAGMIMBA TWKCCCCMAK AACTGAGAAG TOGOGATTAA ATATOGAGTA TIOTCOTGGC CAGGCTTACA TIGMCTCTAG

YWAWCKGAAC AMAMKCYGSW CYTCCWSYGC SKTRRMKRYC GYKSTATRRC WARWKSAKYM

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SEQUENCE DESCRIPTION: SEQ ID NO: 109:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 2539 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

INFORMATION FOR SEQ ID NO: 109:

5

AACTAGAGAG NAAAAAAAA AAAAAAAAAA ATTTAAAAAA CT

TATTITGITIT TITATAACAG GTATIGAAAC AAGTIAACTI GCATICCIAI GTAAGATAGG TATAAAGAAA TOGAAAAAAG TOAAATAAAA AATATOTTOA ATCAGATTTT TTAAAAGGGG AACCTCCTCC ACCCCCTTCC CCTACTCTAG GGGAGAGAGC TGCTAGTGAG ATGACTGTTT AACCICAGGA GGIAACCIIG GGCCCIIICCC IGCIAICCIII ITICICCIIII GGAGGIGCCC

AGGGGCTGAG GGGATCCCCA GTGTTTGGAA CATAAGTCAC TATGCAGACT AATAAACATC

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	(D) TOPOLOGY: linear		TOCAGTCOGC, AGTCCAGCCT GGGGACAGA GCGAGACTCC GTCTCAAAAA AAAAAAAA	1740
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 110:		AAAAAAAA A	1751
S	AGCATGAAGC CGATGGCCOT GOTGGCCAGT ACCOTCCTGG GCCTGGTGCA AAACATGGGT	09	5	
	GCGTTTOGCG GGATCCTGGT GGTGGTCTAC TACGTATTTG CCATCATTGG GATCAACTTG	120	(2) INPORMATION FOR SECTION OF 111.	
2	TTTAGAGGCG TCATTGTGGC TCTTCCTGGA AACAGCAGCC TGGCCCCTGC CAATGGCTCG	01		
2	GOCCCTOTO GAACTTOCA GCACTOGAA TACTOGGCCA ACAACTTOCA TGACTTTGGG	240	3	
	GCTGCCCTGG TCACTCTGTG GAACTTCATG GTGGTGAACA ACTGGCAGGT GTTTCTGGAT	300	00	
15		. 360	(xi) SEQUEN	
	TOTICIONEA TETOGOREAA CETGITICIG GOCCIGATIC TGGAGAACIT CETTCACAAG	420	AAIGITGIGG IGGIAGCAIT 1036TIAAIT CIRAITAIAG AGICICTIGG AGAGCAANGE	9
20	TOGGACCOCC GLAGCCACCT GCAGCCCTT GCTGGGACCC CAGAGGCCAC CTACCAGATG	480 20	_	
}	ACTISTICIAGO TECTISTICAG GGATATTETIG GAGGAGOCCO GGGAGGATGA GCTEACAGAG	540	CAAAGGACCT TGCTAATATC TGTIAAGACG CACCTAGACA ASSERTANT CONTRACTAGACGA	120
	AGACTGAGCC AGCACCGGCA CCTGTGGCTG TCCAGGTGAC GTCCGGGCTG CCATCCCAGC	009	ATGOCCOAN ACTACOPTED ANTOINCES ARGARIAN ANTOINCES	081
25	AGGGCCGCA GAGAGAGAG GCTGGCTAA CACAGGTGCC CATCATGGAA GAGGCGCCA	. 660 25		240
	TOCTETIGGC AGCAGGAG GAAGAGACT TTCCTCTGAC GAACACTAA GCTGGGGACA	720	TOATTCOTTA TOOCTOOCOC TOOCCTTOCA GAACTCCAGT CITTUTCHACT ANAMERA	0 0
30	GRAACCAAOT CCTITIGOSTG TGGCCCAACA ACCATCTACA GAACAGCTGC TGGTGCTTCA	780 30		S
i •	GOGAGOGOCC GIGCOCTCCG CTITICITITA TAGCIGCTIC AGTGAGART COCTCOTCGA	840	CACAACTOT CACAGAAGT CITTITIAGA TAAACGIAGA CITCOTATAS CITAGAAGA	
	CTCCACAGGG ACCTITICAGA CAAAAATGCA AGAAGCAGGG GCCTCCCCTG TCCCCTGCAG	906		0 0
35	CITICOGITICA GCCTTTGCTG CCGGCAGCCC TTGGGGACCA CAGGCCTGAC CAGGCCTGC	35		OF
	ACAGGITAAC CGTGAGTCTG TCTCATCTAT TCACAGCTGG GAATGATACT AATACCTCCG	1020	THAMACTOGN AGAOTOGANA GGCAGACTAC AAOTTACTGA GCACCTCCCT GAGAAAATTG	0 9
40	ATTITIAGOCC AGCACCACAG GOTACCITICC AGITITITCIC TCTITICCATA GCIGIAAGGC	1080 40		220
	CCTITCTGGG AATGGITCTC AITCTCCTTA ATCTATTAIT GGGTCAGITT ICCTGCATGT	1140	ACCITICATED ARACCITICA GIRATAGAIR ARCARGACA GGACTGARAG TGCTCTGAAC	780
	CCCCAGCCTC CCATCACTGC CACCCACTCC CCACAGAGAT OCCCTGCTCA TCCGACTGGG	1200	TTGARACTICA CTOGRGAGCT GAAGGGAGCT GCCATGTCCG ATGARTGCCA ACAGACAGGC	840
45	GCTITGACTC CCACACTOTO TACCCCTCTT GTGTOGAGGC CCTGCTGCCA AAACCTTCAG	1260	CACICITIOS ICAGOCIOCI GACAAATITA AGYOCITOGIA COTGIGGIAS CACITIZATIFIC	
	CANACACTT TCCANATGGA ACTIGTCACT GTCAGGCCTT TACAATCAGC AACAGGAAAA	1320	CICHOTOTT ITICITITIC THITAACIAA GAAIGSGOOG GIIGIRACIC CACHTERACTA	0 9
20	TCTACATIGCT GCTGAGGGTC CTGCCTCATT AAGATGCAAT AAATATGTAA GTACATAAAA	1380 50	ATCCTTAAAT TTAAATACAT ACTTATOTIT GTATTAANCT ATCAATATAT GCAMACATCA	750
	ACAGCAATAG AAGAAACGTA ATGCTTTATT CTCAAATATG ATGTCTACAT AGAAAAGCCA	1440	ATATATICCAC CCACCTAGAT TITTAAGCAGY AAATAAAACA TITTCGGBABA GARTTAAAACA	
	AAATTATTAA GAATAGTAAG AATTCACCCA GCACTTTGGG AGGCCGAGGC GGGTGGATCA	1500	GAATTTTACA CTTAAAAAA AAAAAAAA AAAAAA	080
55	TENEGTCAGG AGATCGAGAC CATCCTGGCT AACAGGGTGA AACCCCGTCT CTACTAAAAA	1560 55		
	TACANANANT TGGCCGGGCG CAGTGGCGGG CGCCTGTGGT CCCAGCTACT GGGGAGGCTG	1620		
9	АОЗСАБВАСА АТООССТВАА СССОЗВАЛОС ОЗАВСТТОСА ОТВАСССВАВ АТТОСОССАС	1680 60	(2) INFORMATION FOR SEQ ID NO: 112:	

WO 98/39448

335

SEQUENCE CHARACTERISTICS

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(A) LENGTH: 1113 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double

(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 112:

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50 25 8 35 30 25 20 15 5 AATOCTITIT TIÖTTATCAG AGATIGIGTA CTATTITIAT TITTAATAAA TGIATCITCC TITITITIAA ITAATAIGIG IOCAIIGITA CAAIGIAIGI IOGGAIGICI TIIGACCCIA AGAAGGAAGC CAAAATAGIT TITICCITIT GAAAGTITIT TAAAAATTAT TICAIGGGIC GACCACGIGA AAGGGAATOC IGGICTAGCI GGCGIGGIAI GITIHTAGGC GAATIICAGC ATTICAAAAAC AATGTGTTCA TCAAAGTAAT TGCTCACATT GTGCAGTACT ATGTTGTACA TITITOTCAA ATOCAACCAT GOATITATOT CIOGATCATC CATACAGAAC CAACAATTIT GTATTTGACA CTCATGCAAA ATAATGTGAA AACATCTAGA TTTAGTAGTT TATTCTGCGC GOTTATITAA GCOGAAGACT ACTITOCOATO CTCCAGGACA TGAAAAGACT GAAGATAATA TCAGTTAAGT AGTIGGTAAC CCTTTTCTAT TITAGTAAAA CTTAATGCAT GTTTACTTTT CTTTTGTTAA AACTGAAGAT TTTGGAAAAT GGTTGTCACT GCTCTTCCAG CCTATGAATA TITIGATOIT TAGAGACACI AGITITICOCO AACITAAGAT TITIACGITAA TITITIACATA TGACAAGCAT TAGTGACAAA GOCAGAAAAG ATTTATCAGC CATGCTAAAA GAGTGAAGAA TOGIATITIT GGIACIGICI TOCITCAGCA GIGCATATIC TITIGCAAAG TICTITGGIT AACTITICACA GICCAGIATC CAACAGGAAC IGIGIGIGIG TIAAGACCGA AGTIACAATA TAMARCAAGC TTACACGAGT GCTCCAATGG TAGACAATGA ATTACTTCGA TIGAGTCTTC TACTACGAAA AGAAGAAAGA GCAAGICTTC TIAGIAAICT TGGCCCAIGI TGTAAGGCG TOTOCTICAG ACOGGATICT GCAATICGAA AGCAGCITOT TAAAAATGAG AAGGGCACCA TITIGAAAGA TOCAGACAAT CICITIGAAC AIGAATIGGG GGCICICAAT AIGGCIGCAI TITIGAGACTA CCAACATAAC TACOTOTTGA AGGTOCTTCA CAGAGAATAT ATTIGCCTTN GCCAGAGGIT TICTIATATI TINAGTAAAT TIAAAGTGGC TATCAGAATA TITATTCTTG ATGTGAAATA ATTTTCACCA ATGTTGCTAA CTTTAATAAA GTATAAAATT TGTAGAATAT 1260 1200 1080 1020 960 900 840 780 720 660 9 8 420 360 180 120

9 SEQUENCE CHARACTERISTICS: (A) LENGTH: 1654 base pairs (B) TYPE: nucleic acid

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CAAAAAAAA AAAAAATAAA NTTCGAGGG GGGC

1654 1620

TAAAATCAGI TIIGIAAATA IGIAAATAIG TCATAAATAA ACAAIGCIIT GACIIATIIC

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INFORMATION FOR SEQ ID NO: 113:

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1313

PCT/US98/04493

WO 98/39448

336

PCT/US98/04493

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

5 TGATGACAAC AACAGTCTTT CATTACAGAC TGAAGGGAAG CATGTCCTTA CTTAAAATAG COTCTACACT TTATTTTAAA AGCTATCCTT TTCTAGTAGT ATTTTATCAT GGCAATGGCA ACAGGGACAG AATACTITCT TICCTICCTT CAAGTACAAG AAGGCITICT CTACCATIIG SEQUENCE DESCRIPTION: SEQ ID NO: 113:

GATTTTTATT CCCAGGATAT GOTGTTCATT TTATGATATT ACGCAGGATG ATGTATTGAG GAATGCAGGA AAAATAATTT AATATCAACC TCAGTTGACA AGGTGCTCAG ATTATTCAAT CAAATTGATA ATATTCTATG TTCTAAAAGT TGGGCTATAC ATAAATTATT AAGAAATATG GAAYCTACTC TGTTCCTTGG CTAGAAAAAA TTATAAACAG GACTTTGTAG TTTGGGAAGC TTCAAAATTA TGCAAGTTAG TAATTACTCA GGGTTAACTA AATTACTTTA ATATGCTGTT AAAAGGTAGT GTGATAGTAT AAGTATCTAA GTGCAGATGA AAGTGTGTTA TATACATCCA TICTITATIC CICTITCITC TGAAGAITAA TGAAGITGAA AAITGAGGIG GATAAATACA TICIGAAIGI TIAGGCAGIG CTAGIAATIT CCICGIAAIG ATICIGITAT TACTITICCIA TOGGGATOOT COTTITIGITA GETTITIGAG ACAACOOTAG ACCTAAACIG IGICACAGAC ACAAATTATT GAATTTTATA AGCTTGTACA CAATATTTAA TTAGTGTGAA AGGAAACAAA TITIGIGATIT TITITITICIT CCGAAGAACT CCIGGITGIT AITGGATITI GIATITIAAT TCTCCCTCTG TCTTAGGAAA AGCCATCTTT AATATAGTTC TTCACCACTG TTGGGGTTGT CCCATTICCT TGAACCICCT GCICTAGCCT TGGCGGAGGG AGAGIGCTAT TIGCTITIGT AGTOTOAGGO CACAGITITOO TICACTAATO GIOCAGOTIG AGIGITOTGI TOTOITOCIG ATGCAGCTT AGCCAGGTAG TCTTAGTGGT GGTGTTTAGG CATAAGATAT GCTGATCATC TCCTTAGATT TCCCTGTTGT AAAAGGGGCA AGAAAAGTAA CTCATCATCT CTAACACACC GCAGCOGANA INGACTGACT TCACATOCTC AGCTTTCTCA GCCTTTTGTT TATTTTTGTTG AGRAATACAT GAACTOCTCT GOCCTCTCTG GTTCTGTTCT TOGCCCAGAG TTTTTTGAAAA OCCITECTOCT COGNACEOTS TEMOTOGOTS AGGNAGATER GAGATOGTER GATEGRAGAG CAGIGGCAGT CCATGGCTIG GITGAAGCIA GAAATITICC TGCCCCTGGT GACCTGGTAA CATAGAATCA CAAATACTGA CATTTCATTA GATGATTATT TTCCTAGAAT CCCCAAAGAG TICTTICATI TITATTITAC CAAGICACAA ATGICTITIT GAIGITITGA GAATIGIICI TTCTGCTACT TTCCCTCCTA TTATAAGGAA ATTTTACAGA TTCTAAAAAT ACCTTAATTT 1560 1500 1440 1380 1200 1080 1140 1020 960 900 840 780 660 600 720 240 360 300 180 120

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<b>NO 98/39448</b>	

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338	(A) LENGTH: 842 base pairs (B) TYPE: nucleic acid (C) STRANDENRESS: double (D) TOPOLOGY: linear	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 115:	GOTOTIGGICO GANOTIGCAT GAOCTIGCCA TOTIGGICCTT AGTICATITICG GITTICGGICG	CICHCCCOSTO TITICCOGGOC TOGGNATITIG CCTCGCACCA TOGGGCCCAA GGGCAAAGTG	GOCACGAGAG GGAAGAAGCA GATATTTGAA GAGAACAGAG AGACTCTGAA GTTCTACCTG	COGNICATAC TOGOGOCCAA TOCCATITAC TOCCTTOTOA COTTOGTOT CTTTTACTCA	.  TOTOCOCCUM MINISTERIOR CONTRACTOR CONTRACTOR DISCOSSION AND CONTRACTOR CON	TACACTOTA TRAGETICIAT GECACIACIA GEOFFICIOTE AGRATIGAGE CONGATIGAT	ACCOMPANYA ACMERATURA POLICARATURA MATERIALEM ACCOMPANYA ACCOMPANY	מסומקישומת שהנובשישו משמבשמתה שומחששת שהנוששת ומומשוההוש	CIGACAGCCA TOGIGCAGGI GCICAGGIOC TICTICICIC ATGICTOGIC CITCLOGGIT	CHOSCIPCIAGO COCOGOCOCOT TRACTICOTO TOCOGOANIO TOCOGOCOCO CHOSTICACI	GCAGACAGTG GCACCCCAGC ACCAGAGCAC AATTGAGAAAC GGCAGACGCCG ACAGGAAGCGG	COOCHANTSA AGGOSTIANA GCCATTONCO TIONGOCCAC AGGCCACIUG CCCTUGGGTUG	CICTOTCAGO GIGCACAGCC CCICATOCCT GOAGCAATGA GAGICTAGTC CAGAGACCAA	ANGCRETCIG AGGIRITGGS TRIALITRIR CICTATAGGS TUGITGANTA ANTUGULTAD	ARIUTGARA AMAMAMAA AMAMACING AGGGGGCCC GGFACCHARF TICHNINANA	A		(2) INFORMATION FOR SEQ ID NO: 116:	7)		(C) STRANDELNESS: GOUDLE (D) TOPOLOGY: Lineax	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 116:	GOCACCAAGC GGCGGCAACG GTGGCGGGGG CGCCCCCCGG CGGGAGCCGT TCCCTTTCCC	STCGGGGAGC GCGGGGTCGG GGCCCAGGGG ACCCCGGGCC ACGGAGAGGG GGAAGAGGAT	GGATTIGCCCG GCCCTCCCCC CCGGATGGAA GAAGGAGGAA GTGATCCGAA AATCTGGGCT	ANCTICACIO ANGRICIATI CATACTACTA CAGTICCAAGT GOTAAGAAGT TCAGAAGCAA	GCCICAGITIG GCAAGGTACC TGGGAAATAC TGTTGATCTC AGCAGTTTTG ACTTCAGAAC	
		9		01			60 15	120	180	240	300	360 25	420	480	540	009	660 35	720	780	940	006	960 45	1020	1080	1140	1171	55		09	•
337		(2) INFORMATION FOR SEQ ID NO: 114:	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1171 base pairs	(B) TYPE: nucleic acid (C) STRANDEDRESS: double	(D) TOPOLOGY: linear	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 114:	GCCAAACTTT CCCCCAANGC TTCGAAACTT GCAAGOCGAA ACCTTGAATC GTJAAAAGTT	GOSTITOCONE GOCOCCETGO COCGAAGAAG CGCAATTGOC GTTCCGCGAA COTTGGCCCT	CAACGGCTCG GCAGCCAGGC ATGTCCTGCA CCCAGGACAG CGGCCCTGGG CTACAAGGAC	CTGGACCTCA TCTTCCTGGG CCGACCTGGG CGGGGTAAGG GGNATTTCA GACTGTGAAG	GACCICCIUC TOGACTOCCT GTTGGACTTC TTACCCGAGG GGGTGAACAA AGAGAAGATC	ACACCACTCA COCTCAAGGA AGCTTAATGTO CAGAAAATGG TTAAAGTGTG CAATGACTCT	GACCGATGGA GTCTTATATC CCTGTCAAAC AACAGTGGCA AAANGTGGA ACTGAAATTT	GIGGATICCC TCCGCAGGCA GITIGAAITC AGIGIACAIT CITITICAAT CAAAITACAC	TCTCTTCTGC TCTTTTATGA ATGTTCAGAG AACCCAATGA CTGAGACATT TCACCCCACA	ATAATCOGGG AGAGCGTCTA TGGCGATTTC CAGGAAGCT TTGATCACCT TTGTAACAAG	ATCATTGCCA CCAGGAACCC AGAGGAAATC CGAGGGGAG GCCTGCTTAA GTACTGCAAC	CTCTTGGTGA GGGCCTTTAG GCCCCCCTCT GATGABATCA AGACCCTTCA AGGTATATG	TOTTCCAGOT ITTTCATICGA CITCTCAGAC AITTGGAGAGC AGCAGAGAAA ACTGGAGTCC	TATTTGCAGA ACCACTITIST GCGAATTGCA AGACCGCAAG TATGAGTATC TCATGACCCT	TCATGGAGTG GTAAATGAGA GCACAGTOTG CCTCATGGGA CATGAAAGAA GACAGACTTT	AAACCTTATC ACCATGCTGG CTATCCGGGT GTTAGCTGAC CAAAATGTCA TTCCTAATGT	GOCTABATOTO ACTIGCTATT ACCAGOCAGO COCCTATOTA GCAGATOCCA ACTITAGGAA	THACHACHTI GCACAGTITC AGCCAGTAIT CACGTGCCAG CAACAGACCT ACTCCACTIG	GCTACCCTGC ANTRAGAAT CATTTAAAAA TGTCCTGTGG GGAAGCCATT TCAGACAAGA	CAGGAGAGA, AAAANAAAA AAAAAAAAA A			(2) INFORMATION FOR SEQ ID NO: 115:	(1) SEQUENCE CHARACTERISTICS:

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CACACAAGCT CTGCGCCAAT CACAGGGCAA GTCTCCGCTG CTGTGGAAAA GAACCTGCTG CTITHOTIGCA TCAGATOTAA CAGAACAAAT TATAAAAACC ATGGAACTAC CCAAAGGICT TTTGGCTTAA CACATCTCAA CCCCTCTGCA AAGCTTTTAT TGTCACAGAT GAAGACTCAG CTTTATTGAA AGGGGACACC TGTACATTCT TCCATCGTCA CTGTAAAGAC AAATAAATGA TAGCACTTAC GTAAAACATT TGTTTCCCCC ACAGTTTTAA TAAGAACAGA TCAGGAATTC TCAAGCAGGA CCCTAAGATG AAGCTGAGCT TTTGATGCCA GGTGCAATCT ACTGGAAATG ATTATICAAA AAATCAIGIT TATITIGAGI CCIAGGACII AAAATTAGIC TITIGIAATA TAGAGETTIT TAATAGEACT AACCAATIGEE TYTTTAGATG TATTITTIGAT GTATATATET ATTGAACAAA AATGTTTCCA CTGGCTTTTG CCTGTAAGAA AAAAAATGTA CCCGAGCACA GCCTAAGAAT ATGATCAGGT ATCTTGTCGC GAGCTGCTGA TACAGAAGAG GAAACAGAAG AGCGAGTACA GCAAGTACGC AAGAAATTGG AAGAAGCACT GATGGCAGAC TCAAGGAGTT GGTCCAGTAG CAATGATGAG ACCCTTTTAT CIGCIGTIGC COMPRIMINGS MATICATINE ATTENDATION ACTICIDATICS COGGOCCITA GOTTICACAT TAAATAAATT TCCCAGTTAA AGATTATTOT GACTTCACTG TATATAAACA TATTTTTATA TTATATICCA CAGAAAAAAA AAAAAAAAAA MMSTYGARRR GSROCMCRSM AYMMARWWCC TOTOGTAATT TITTTTCCCT GGGAGGTGGG INFORMATION FOR SEQ ID NO: 117: E (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 117: AGGAAGATAG COCATATATT TOCAGTATGA ACTATTGCCT CTOGGACOTT GIGCITICAC SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double g (A) LENGTH: 952 base pairs TOPOLOGY: linear AACTITICGAC CGACTITICCC CAAGAGAAAA TITCCTAGGAA CAGAATTICT AAGGATTICT GGCTTAAATA TCACCTAGCC ATGGATATTG AAATGGACAG TGGAGATGAA 1200 1380 1260 1140 1080 1020 1440 1320 1620 1560 1500 960 90 840 780 720 66 60 1640

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(2) INFORMATION FOR SEQ ID NO: 118:

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SEQUENCE CHARACTERISTICS:

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1256 base pairs TOPOLOGY: linear

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25 20 5 မ 2 S TOOTGACOTT OTHGATGAGE AACAAAAAA TGAAATAAGT TOTTGGAAAT TAAGOCATTT CATCTITIGGS GTAAAGAGIT AAGTGTCCAA AGGTTGTCAC AGTTCATGAG GTCAGAGGGA CTRAGGRTAG TETTTTRAAAA CTCCCAAAGA AAATCTGCTC TCCTTTCTGA TCTRAAAAACT CTITITIOGIO AACTITAGIG GACTICIGIG AGATIGIAGI IGIACITIGI AICICIAAAI ATTITICCIOC CAGATACCITI TATICAAAATT ATTOOCCITCA TGAGAGCIGA AGTAAGICAG ATTITAATIT GCTATITITI TCAATGITCT AGGTATCITT AAATTIGITA TIGIGGAATC ATTCCCTTTT TTGAAACAGG AAAAAAAATT ATTTTTTGTT CAGTAAAAAT GGTAGAGAAT TAARTICTCC AGTAGACAAT GCIOGGTAAG GGAGGGGGTA GGOCTGGGTT ATTAAGATAC GCTNGCCTGG CACCTGGACT CTGCCCATCC ACAGCTGACA GATTCCAACA GAAGTGTATT TATCTCARTA ARACCCCACT GGRACTCCAA ARAAAAAAAA AAAAAAAAGA NN ATTYCAGAAA CATTOGGGGA AGGGAAAATT GGCTTYCTCT TAATTGGCAG ATGTTCCAGT AGOCTGCTGT ATTITACATT GGTTGTGGGG GAAGGGGAGC CTGGAGAAAA CAAAGTCACT COGAGICTOC TOAGITTATA AGGITCCAAA AATATOGIAA AATCITOGIT TITGITAAIT GOGSSOGOG GOCTCTOTT TIGTTGOGAT GTGTTATGTT GTATGTAGGC ATATATGGAC TOCAATOTOO OTAGOOACAA GGGACCAGTT COACTGAGAA GTGAACAGTG GGAACTCAAA 540 80 420 300 180 120 952 900 840 780 720 660 600 360

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CARTTITICAA ACAACCOGTA ACCCAAAGTC ACAAATCATC CTAGTAATAA AGTGAAATCA

CAATCAAAAT AAGGGTAAAC CAGACTTGAA ATACAACATT GCCAATTAGA CAAACAGCAT TOGANAGATO ATOCCTAGTA AATTACAGAA GAACAAACAG AGACTGCGAA ACGATCCTCT

GACCCACAAC GAATGAATGA ACAGCCACGT CAGCTTTTCT GGGAGAAGAG GCTACAAGG

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480 420

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GACGTCATAG GTAAACAGGC TCTGTATCCG TGGCAGCGGC CGTGGCAGGC TGGCTGGGTA

COGCTOTOG CTGACCCAGG AGAAGCTOCC TOTCTACATC ACCTIGGGCT GCAGCGCGCT 120 8

ATCITTOTAC CCCCAGCATC TAGCAGTOTT GOCATOTAGT AGGCACTCAA GAAATGTGTG GOOGCOGOG GACCGGCAGC TOWACTATOT GCTCTTCAGG GCGGGCACCG TGTTGCATTC 180 240

ATGACACAC TECTECTGAC TOCCACTOTE ACTECTTEAG AGCAGAACTE CTETAGGGAA CCTUGATUGG AAACAGCCAT GGCCAAGGAC ATCCTUGGTU AAGCAGGGCT ACACTTUGAT

TIGANIGANC GAIGCCIGIG ACANGCANGC GGACTITATI CITICCIGAC CCITOCICCI

8

TGAATTTAGN AAACACTTTG GAAAACTCAT AACCTCATCA GAAACTGCCT TIAGCCACAC

GAACTGAACA AGCTGAGGGT GTTGGACCCA GAGGTTACCC AGCAGACCAT AGAGCTGAAG

420 360 8

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3 S. v 2 2 ន 22 20 33 **4** 45 S 25 8 540 80 99 420 480 180 240 38 360 540 909 99 720 780 840 900 960 020 0801 1140 1200 1256 ŝ 120 CACTICATIC TAAACACAGC TICICICCIG AGIGIACIAA TICCCAAAAT GCCACAACIA CARGERGITC GGAICTITIGG AATTAATAAG TATIGAAATG TITIGAAACT GAAAAAAA ACAATTICIC CAATAATICT ATATITICIG GCAAGITICT ATAGGAAGIA TGAICCAACI TICAGIGITIC CIGITGCITIG GACTITIAACA AATATIATAC AIAAICIGGG GAIGIACGIA TITITICATG CAGTGAAAGG AACACCTITC GAAACTCCTG ACCAGGGTAA AGCAAGGCTC CHARTCAIT GOGAACAACT GGACTATGGA GTACAGTTTA CATCTTCACG GAAGTTTTC GAAGAGTGCA AAGACTTTGT GGACAAAATT GGCCAGTTTC AGAAAATAGT TGGTGGTTTA GCTCCGAACT TGCTCAAATC TATAGCAAAG CAGAGAGAAG CTCAACAGCA GCAACTTCAA OCCCTAATAG CAGAAAAGAA AATGCAGCTA GAAAGGTATC GGGTTGAATA TGAAGCTTTG ICTAAAGTAG AAGCAGAACA AAATGAAITT AITGACCAAT ITAITITITCA GAAATGAACT GAAAATTITCG CTTTTATAGT AGGAAGGCAA AACAAAAAA AGCCTCTCAA AACCAAAAAA ACCICTOTAG CATICCAGCO GCITGACCAA TGACCIAIGI CACAAGAGGI GGCGTGIAAG GAATOCAGCC CCCTGAAGAC AGCACTACAA GTCTGGGGGA GCCAGTTTTA ACATCAGTGC ACCAGITITOT AAACITITCAT CCITITITIT GIAAAITCAC AAACCITITOG AAGGAAAGC AATAAAITIT TOTITICAAA TOOCTIGAIG TACCITITIT CCTOTIGCIC IIGAAATAIG ITTAACTCCT CATGAGAA CCCTGGATTC TCTATCCCCT AGTCCACAAA ACAAACCAGG GOCCOTAGCA GOCGGGCTGG TOCTGCTGCG AGCGGGGGG COGGAGTGGG GCGCCGCAT GHACCTICCA CATTGAGIAT TCAGAAAGAA GIGAICTGAA CICTGACCAT ICITTAITGAA TACATTAAGT CAAATATAAG AGTCTGACTA CTTGACACAC TGGCTGGAGC AAACATGAAC STROGAGITIC CCCACAGIGA AGTGAAITCCA AAIACCCGTG ICAIGAACAG CCGGGGIAIG TOSCIGACAT AIGCATIGGG AGTIGGCTIG CTICATAITG TCTIACICAG CATICCCITC ATTGACCTTG TTGATCAACT TGCAAAAGAA GCAGAAAATG AAAAGATGAA GCCCATCGGT GTTGGAACTA CAGTOSTICAG CACCTACTT TINITITICAT CACACACTG AGTCAGACAG TACCAC ACAGCIGCIG CIGGIGGCCC TGCAGTGTAC GITCTCACCT CITATGCTTA SEQUENCE DESCRIPTION: SEQ ID NO: 119: LENGTH: 1143 base pairs TYPE: nucleic acid STRANDEDNESS: double SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 119: TOPOLOGY: linear 3 X; Ξ 3

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780 840 8 140 8 249 8 9 720 ද 120 8 8 AGAGGGGCTT TATAAGCAGG CTGGGCAGGC CCAGCTTATA AGTTAAAGGG CATCACAGTG TITACACCTA CICAATITICI TATAAGGAAG GAGTGGITAG TAAACIGCAC TGITTICIISTG TITGAAGIGC AGAITICCAI TAAAIGAIGC CICIGITTAA TACACCIGGI ACAITITCIGA AGGOTGTAGT AGATABATTC AAGGABATAA GAGATTTGTA AGABACTAGG ACCAGCTTAA CITATAATGA ATGGGCAITIG TGITAAGAAA AGAACAITITC CAGTCAITICA GCTGTGGTTA TITIAAAGCAG ACTTACATGT AAACCGGAAT CCTCTCTATA CAAGTITATT AAAGATTATT TCTTCAGTTT TACAGAGATC CTGGGTGTTT GGCTGACCTA CAGATACAGG AACCAGAAAG ACCCCCCCCCC RANTCCTAGT GCATTCCTTT GATGAGAAAA CAAGGAAGAT TTCCTTTTCGT ATTATGATET TOTTCACTIT CTGTAAITIT CTGTTAAGCT CCATTTGCCA GTTTAAGGAA ATAATGIGAA ATGAGAAGTA TTTACATTGG AGGCCAATG GCTGGTCCTT CAAGTGCTGT TITIATTACCG TARABABAR ABABABARA ABABABARA BARABABARA ABARABANA CAGGCCCCCG CCCCCCACCC ACTICTGCGT TGCTGCCCCG CCTGGGCCRG GCCCCAAAGG ACCCAANIGA CACCTIOTICTO GCTAGCTOTO TTAAAAGTGA CCACTCGTGC TCGCCATGTG CICCAAICAT AGGAGAAIAT GCIGGAGAGG TITIGAGAIT IGIIGGIGGC AITIGGCCIGI CAAGGACAAA GCAGCTGTCA GGGAACCTCC GCGGAGTCG AATTTTACGTG CAGCTGCGG CAACCACAGG TTCCAAGATG GTTTGCGGGG GCTTCGCGTG TTCCAAGAAC TGCCTGTGCG COCTONACCT GOTTTACACC TTGGTTAGTC TGCTGCTAAT TGGAATTTGCT GCGTGGGGCA TIGGCTICGO GCIGATITICC AGICTCCGAG IGGICGGGG GGICATIGCA GIGGGCAICT TCTTGTTCCT GATTGCTTTA GIGGSTCTGA TTGGAGCTGT AAAACATCAT CAGGTGTTGC TATTIVITITA TAIGAITAIT CIGITACITG TATITAITGT TCAGITITICT GTAICITIGC CITICITITACC CCIGAACCAG GAGCAACAGG GICAGCITICT GGAGGITIGGT TGGAACAATA CGAAGTGTTA COSCAACTIC TCGAAATGAC ATCCAGAGAA ATCTAAACTIG CTGTGGGTTC (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 120: (A) LENGTH: 1782 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDMESS: double
(D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 120:

TACTOGAAAA AGAGTOGRAA TITATTAAAA TCAGAAAGTA TGAGATCCTG TTATGTTAAG CTTAGCATTT TTACCTGCAG AAAAACTTIG TAIGGTACCA CIGIGITGGT TAIAIGGIGA GTAGCAAAAA GATATTTGAT TATCTTAAAA ATTGTTAAAT ACCGTTTTCA TGAAAGTTCT AATOGAACGA GTTTTGAGTA ATCAGGAAGT ATATCTATAT GATCTTGATA TIGTTTTATA GAGAAGTOGT TICATGAAAT GITCTAATGT ATAATAACAT TTACCTICAG CCTCCATCAG AMATAGITAT GYCYTMOGAM ATTGIGGTIT MATTTTTGAC TITTRCAGGT AMGTGCAMAG TOTTAGTATA AAAATGATAA TIWACTKGTA GICTITIAIG AIWACACCAA IGIAITICIAG GGAAATCCAA ATTCCCAATT TTTTTTGGTC TTTTTAGGAA AGATGTGTTG TGGTAAAAAG ATCTGAACGT ACATCTCACT GOTATAATTA TATGTAGCAC TOTOCTOTOT AGATAGTTCC GAAATTGAAA TCGTATTGTG TGGCTCTGTA TATTCTGTTA AAAAATTAAA GGACAGAAAC CAGTATTOTA ACAGCAACTT GTYAAACCTA AGCATATTTG AATATGATCT CCCATAATTT ATAATTIGAA GICIAAAAGA CIGCATTITT AAACAAGITA GIATIAAIGC GIIGGCCCAC TTOGATAGGA ATGGGGCTGA TGGGCTTCAT CGTTTATAAA ATCCGGGCTG GTTGGCTGCA GATTTGTGGT GCGTTCTGAG CCGTCTGTCC TGCGCCAAGA TGCTTCAAAG CTTTCTTTGT GTATGCATGT TTGAATTAAA AGAAAGTAAT GG TATTATTADA AACATATOGA TCCCCATGAA GCCCTACTAC ACCAAAGTTT ACCAGGAGAT OCCITATIONG CATACTOTICA GENGALINATE TOTOLITOCHG COCATOCHAE CAGAATOTICA GATCTOTOAG ATGCACTGCT ACCTGGTACT GCTTTCAGTG TGTTCCCCCT CAGCCCTCCG GTACATGTGA AAGAAAACGT CAGTCTGCCT GTAAATTTCA GCAAGCCGTG TTAGAIGGGG AGCGTGGAAC GTCACTGTAC ACTIGTATAA GTACCGTTTA CTTCATGGCA TGAATAAATG AAGTAAGGCT TIGAAAAGCTT CAGCGCCTGC TCCTGGTCAT CACAACCAGA (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 121: (A) LENGTH: 610 base pairs TYPE: nucleic acid STRANDEDNESS: double TITACITICA CTGATAAAAG 1140 1080 1320 1260 1200 1740 1500 1440 1380 1680 1620 1560 180 120 240 360 300 420 S 480 23 20 5 8 5 8 2 6 ઝ S 8 TTCATCCCTT CTACCCATCA CAACATACAA AGCAACTACA CCTGGATTTT TCCAAACAAC CAGGOTTCCA GGACATAGIC TGAGGCAAGA TGGAGGGTAT GAGGGGCCTT CACACTTCAC GENGETEGEE ATETTEGEEN ACATGETGGG COTOTOGETE TECTTOCTTG TEGTTETETA GOTACOCCTG CAGGTACCGG TCCCGAAATTC CGGGTCGCCC ACGCGTCNGG CCACGCGTCC 2 CCAGTATAGG GGCTIGCTTT TCTACTCCCT CCCCCCAATA TAAAAATATA GACTTTTTAA TCACTACOTO OCCOTCAACA ATCCCAAGAA OCAGGAATGA AAGTGGCGCT TTCTCCGCCC ACCCACOCOT CCGSCCACOC GTCGGAGCCG AGCCGGACTG GTCAGGATGA TCACGGACGT TITTATITICC TCAGAGICIT CCTTAAICCT AIGGAACAAG AAGCIGCCAC IGAATAGGGC 5 AAAAAAAAA AAAAANTTOG NOOGGGGCC GGTACCCATC CCCCTA ACCIGCAMATT COTOGTOCAG CTGTTCGCCG AGGAGTGGGG CCAGTACGTG GACTTGCCCA COGGCTACGA GCTGCTCATC CAGAAGTTCC TCAGCCTGTA CGGCGACCAG CAGICCCOGI GCICTICIGI TICICAGICI ICGCGCGACC CICGICGGIG CCACACGGGG AGEOGRAPH CETTEGORTE CECAACAGEG GEGETGGGGG GEGGGGGGC CEGTEGGGCA TOTACCOOTC COGAAATTCC COGOTCGACC CACOTCOTCS GOOGAACATG GCGGCTKCGG INFORMATION FOR SEQ ID NO: 122: INFORMATION FOR SEQ ID NO: 123: Ξ E (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 122: ž SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 123: 0 (B) TYPE: nucleic acid (A) LENGTH: 526 base pairs (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 2081 base pairs TOPOLOGY: linear STRANDEDNESS: double

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ATCGACATGC

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(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 121:

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SEQUENCE CHARACTERISTICS:

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GUICGOTOGO ACCIGGGAAT TIACIGTATI CATIGICOGG CACIGICCAC TOTOGCCITI

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AAAAAAAAAA

CCAGGCACTC ACTIGIATIC TACTGCTCAA TAAACGTTTA TIAAACTTGA AAAAAAAAA

8 919

CTGAOCCACC CATCATTOTO AAATAATTAC CTCAGTTOTA CAGGACTTOG TGATCAGGAT

960 900

TICTCIACAT GITTITITCT TICCGITGCT GAAAAATAIT IGAAACIIGI GGICICIGAA

GCAAACACTA TCTOGAAAAG TACCTTATTG ATAGTGGAAT TATATATTTT TACTCTATGT

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сствалсесь святаваеть ввевсловае свалоссвое всевствалае GAAAGGAATT GCCCTTACTA CTTTGGGTTT GGTTTGCCCT TGGCTTTTCT CACAGCAATG TIGGIGGICT TOTTANGCAA CAGACTOTTO CACAAGACAG TOTACOTGCA GTOGGOOOTG AGCAGCICTA CITICIOCAGA GAAGITICCCI TCACCGCATC CGTCGCCTGC CAAACTGAAG OCTACTICIAG GICACTICAGT TGCCTGCCAT CCAAAGGGA TGGCGGGAT TGGAAGAAGC TOTOGCAGET CITITICECTO TICACETCCE GCCTGCCAGG GNAGGCAGGA CCCGCTCTGC CAAGGGCCCT CTGCGTATTC CCTTCTCTCT GAGGAATTGA AATTTTTGTC TCTGGTGCAC IGCTTTGAAT ATCGTTGGTT CAATAAAGGA AITGAAATGC ACCAGCGGTT GTCTAACATA CAGICCICAT ATATTAICAG TOGCTOCCTT TICTCTAICC TCTTTCCTTT ATTCATTAIC AGCOCCAATO AAGCAAAGAC CCCTGGCAAA GCRTATCTCT TCCAGTTGCG CCTCTTCTCC GIAAGGCAGA AIGIICCCIG ACACCAGIGI GIGGAITITIT AACAICACCG IGAGICIGAA IGROGATICET TEATGATGAG AGATTITGGGG ACACTICTET CTECTIGITG TAGITGATAG ITTIGGTGGTG AAGAGATGGC TGACAGTGTC AAAACCTTTC TCCAGACCT TGCCAGAGA ATCAAAGACT CCATCTGGGG TATTTGTACC ATCTCAAAGC TAGATGCTCG AATCCAGCAA AAGAGAGAG AGCAGCGTCG AAGAAGGGCA AGTAGTGTCT TGGCACAGAG AAGAGGCCAG agtatagagc ggaagcaaga gagtgagcca cgtattgtta gtagaatttt ccagtgttgt GCTTGGAATG GTGGAGTGTT CTGGTTCAGT CTCCTCTTGT TITTATCGAGT ATTTATTCCT GTGCTTCAGT CGGTAACAGC CCGAATTATC GGTGACCCAT CACTACATGG AGATGTTTGG GIGCTITACCA AAGIGGIGAA IGCCATITIGG ITITCAGGATA TAGCIGACCI GGCATITIGAG GGAAGCCTCA CCCATTCCCT AGTGTCAGCA AAATAATTGC TGACATGCTC TGTTTGTGAG TCTCTTTCCC ATCCATCTTG TCGGTCAGCT GGTTAGTCTC CTGCATATGT CCCTTCTCTA CTCACTGTAC ICGIGGCIGG AATICITICCT CACGICAAIT TICAGIGCIC ITTGGGIGCT CCCCTIGITI SEQUENCE DESCRIPTION: SEQ ID NO: 124: 346 TICAACCITY TECHGCAGGC TETTITICETC AFTCAGGGAA (A) LENGTH: 1717 base pairs.
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 124: TOPOLOGY: linear 9 Œ 3 DOCCOGCOGA GTATCAGGGA 3 2 2 2 23 8 33 6 5 20 55 8 2040 380 420 480 \$ 1740 1800 1860 1920 1980 80 99 720 780 1440 1560 1620 1680 2081 8 8 960 020 080 1140 1200 1260 1320 1380 1500 ATTCAAATTA TITTGACCTC CACAAGATGG CTCTATGAGT ATTCTTCCAG GGATTCTAAT ICICITICGAI ÁAAITITAITI ICAITAAATA CITRITAGAG GGIITIGAAA TGITITICAA ATATOTGAAA TOTGAAACTG CTGTCTTTTA TATTAAAGTA ATTAAAGAAA ATGTATTGTG AGGOTTICGO GGTRAGOGAG CGCTGCAAGG TGCGCCTCGT GCCGTTGCAG ATCCAGCTCA CHACCCIGGG AAAICTTACA CCTTCAAGCA CIGIGITITIT CIGCIGIGAT AIGCAGGAAA GOTTCAGACC AGCCATCAAG TATTTTGGGG ATATTATTAG CGTGGGACAG AGATTGTTGC AAGGGGCCCG GATTTTAGGA ATTCCTGTTA TTGTAACAGA ACAATACCCT AAAGGTCTTG GGAGCACCOT TCAAGAAATT GATTTAACAG GTGTAAAACT GGTACTTCCA AAGACCAAGT ITTCAATGGT ATTACCAGAA GTAGAAGCGG CATTAGCAGA GATTCCCGGA GTCAGGAGTG ITICIATTATT TGGAGTAGAA ACTCATGTGT GCATCCAACA AACTGCCCTG GAGCTAGTTG GCCGAGGAGT CGAGGTTCAC ATTGTTGCTG ATGCCACCTC ATCAAGAAGC ATGATGGACA GGATOTITISC CCTCGAGGGT CTCGCTCRAR CCGGGATCAT AGTGACCACG AGTGAGGCTG THAAGGCGAG TECTECAGAG TEGGGTETGE TITCCAAAGT ATAGGACATT TGAAGAACTG CCOSTOCTGC TTACCTTCCT TTTTTGTTAA TGTGCTTTTA TTTATTAAAA AAAATTACAA CTTOTGAAGT TTTCTTAAAT TGTTCACTTT AAAGAAAATG ACGTACCAAC AATGATTTGG CITITATATT ACTORAAGAT GITATAANGT TAANGIGGAT GIAGIGGITT TACITITACAG GITGAATAAT TACATATCIT ICTIGACIAI ACIGAÑIICT TAÑITIGGIC ACTATIACIA AATCTCTGTT AATATTCTCT CTTTTAACTG AAAAGGGATG GGATAGAAGG GTTTGCAATG ACCAACATUT TGAATATATA TTCTAGTGTC CACAAGATTT AGCAAAAGA TAAAGCTTGG GIGGAATATC ATTITAAAAT GITCATGITC IGITCIAIAT ITTCITCACC TACTCICCAA ataticidar gcaaaagic icagiaaiga titogiagia itaaititigi goicaficit INCHECTICA GCIEGIAGCI GAIAAGGACC AICCAAAAIT CAAGGAAAIT CAGAAICIAA STATECTACT CACTEGIGAA GGACAGTCAG GTGAAGGACT GTAAGCCCAC ACAAGCTCTT CITATCICTA CTAGAATTAA AATGITAAGI CAAAAACGGC ICCITTITITG CGCCICCTAG IGAAACITIAA CCAGCIAGAC CAITITGAGIA CCAGCAITITA GITACAAACG ICAAAGGCIT IGAAGATGCC TGTTTTGTCT CTACTGTGA CTCTGATCGT ATCTTTCCAA AGTGCAGACT ATTCATTGGA ATAAGATTAT TGCATATGAA TTTACCCACA GGACTCTGAA TCATGTTACC CACTCCCCTC ACAATOTIGT CCACTTAGIG AGITGCATIG ATCTAICCGT ACCAAAIGAT CCATATTATT GGTGGAGGGC TGTTTFAACA TCTTTGAAGT ATGGCTTGCT GAATATCTTT ATTIAITIAA GGINATAAAA TCTTGACAIT TATAATCIIT C 3 2 2 25 35 6 20 55 8 ឧ 3 5

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1680 1717 480 420 360 8 240 180 120 600 540 804 780 720 25 6 35 မ 20 8 ន 8 5 5 5 GOGTTATTTT G GOCCATOGAG GCATTICTIT CAGOGAAATG OTCCATNATT TCAGCCAGAA GGCATTOCAT GIGGGCAGGI CAGIGATIGIC AGCAGIGGAG TGATICCCAG CACAGCGGCI TCTGGGAAGA GCTGGGGTQA ACTITATITT AGCCCTTCCC TIGTTGCTCT TATGGAAGAA CAGAGGAAGG GRAGOGICIO GRATIGOGOCT OCCCCIGANO OCCCIGANON GRAGIACCIT GCCAGCAICI THIGHTACCA CAATTAGAAC TOOCAGAGCA CTGAAAGAAA AGACTITTOCT TCCCGAAGAT CTTCAAGGAG GGGGCAAAAN GACCTTTAAG TTTTTTAGGTT TAACACAGGG AACCCNCAAA TANGTTANGT CONSCRIPT TOTOGCCCNG CTCTOTOTTA TINNGGGCCC TTGGCGANGA CCAGAAGGAA CTOGNGAAAG CGAACAGGAA CCGTTCAGGT CCTGAAGCGG TCAGGCCGAG GATIGETTIBE AGGEATTIGGT TGACTIGGTTA TACAAGGTIGG AGCCACAGCT GGCTGAGGAC AAGICIGIGG ASTCAGAAAC TIGACAATIT CCIAGGAGAA GICAGAGACA AAIGGGATAC IGITIGIGG AAACTTCAGC TTTCTAAGCA TAAGGAGTTT CAGAAGACTC TTGGTGGCAA GCAGCCTGTG TCTCTCATTG ACACCCATAA GGAATTCATG AAGAAAGTAG AAGAAAAGCG AGTGGACGTT CCTTANANCA AGCOGNAGTO TITICGAGACA CAGTICCACAT GCTGTTGGAG GCACTCGCTG GGACACTGTC TGTAAACTCT CTGTTTCCAA ACAAAGCCGG CTTGAGCAGG CAGCCCCGTGC NGGCACGAGG AGAGTCACCT GGACTCAGAA CTAGAGATAT CCAATGACCC AGACAAAATT 5 AAGCAGAGCA AACGCTTOGC TTTCGGGGAG CACTTCCTGG ATGACACAGA GGCCCTGCAG AGCTGATTGA GAATAGTCGA GATGACACCA CTTGGGTAAA AGGACAGCTC CAGGAACTGA INFORMATION FOR SEQ ID NO: 127: (X Ξ SEQUENCE CHARACTERISTICS: ACCOCIOGACE TIGACETECT CATGAACETE ATGGATGCAC ACAAGGTTTT AGCGGCAGCA CAAGTTIGGAG GAAGCCCTIGC TCTTTTCGGG TCAGTTICATG SEQUENCE DESCRIPTION: SEQ ID NO: 127: (C) STRANDEINESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 3752 base pairs TOGCTTTCTG 431 420 240 120 660 600 540 480 420 360 300 240 180 120 180 8

GOCACAGECE AGGOCETTGA AGCEAGETGG CECTGGAGAG GGGCTGETGT GECAGETTGG (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 126:

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STRANDEDNESS: double TYPE: nucleic acid

TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 125:

(1) SEQUENCE CHARACTERISTICS:

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TIGCICCIGG TCACTCCCTT TATAGCCATT ACTGICTIGT

TICTIGIAAC TCAGGITAGG

1620 1560 1500 1440

TITITOGICIC ICTIGCICCA CIGCAAAAAA AAAAAAA

ATTITICITICT TITICIGIGA AACACATACA TIOGATAIGG GAGGTAAAGG AGTGICCCAG CTACCACCCT TOGATGAATG GATTITGTAA TICTAGCTGT TOTATTITGT GAATTIGITA ATTIGGAACAC TITIGITITITIC TECETIGICE ATTITACECTT CEACETTICE ATECTICECT AGGACCACAG GITITICIOC AGCIATITIC TAGCATITOC CAGICCCIOI OCCIGGACIO

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SEQUENCE DESCRIPTION: SEQ ID NO: 125:

CCACGCGTCC GGTCACTATG TAGTGGAGGG GCAGACACCC TCCCGCAAAT TCTGGAAGGT

TETTAGTETE GACTAGGGCA GTAGECCCAG GACTECTAGT CGCCGGCTTC AGGTCACTGC

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COSCIDANCO GNACIOCCOT COCCNIGITY GACIOCITOS TRACCISARAS OCTOGIOCAN

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INFORMATION FOR SEQ ID NO:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 431 base pairs

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NINTITIGGN ANACATAATI TGAATAAAAT AATITITAAT GGATINIGNA AAAAAAAAA

CCGGCAAATG TGGTTCTGAA ATGGTATGAA AACTTTCAAA GACGACTAGC ACAGAACCCT OCTICATICAT TIGGIGICIC ICAGGCCCAG ATGACACCAA GCCCATCIGA AATGITCATI TOCTOAGTTG ACTOATTCAC TOAGTTCACA CAAAAGATGT TGGACAATTT CTACAATTTT ATTICAGIGG AATTATTAGA CAGTAIGGCI CAGCAGACIC CIGTAGGIAA IGCIGCIGIA 8

AGCCAACATC CTTTTGGAGC CATGAATATT GTCCGAACTC CATCTGTTGC TCAGATTGGA OTCACGANTO GGAAGCCAAG TGCCATCTTC AAAATTTCAG GTCTTAAATC TGGAGAAGGA TOTOTOTACT TITOTIANCO TGATTCAAAT GGAATGCCAG TATGGCAACT COTAGGATTT ATCAACCATG TIGIOGITITI TAIGCIGGGA ACAAICCCAT TICCIGAGGG AAIGGGAGGA ACAGCTOCAC AGCAAGTOOC AGAGGATAAA TTTOTTTTTTO ACTTACCTGA TTATGAAAGT

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

348

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840 8 960 1020 1080 1140 1200 1260 1320 1380 1440 1500 1560 1620 1680 1740 1800 1860 1920 1980 2040 3100 2160 2220 2280 2340 2400 2460 2520 2580 AATCCTCCTC CATACATICG OFGTATATIT AFTCTGAACG GGAGAAGTTA TATTGTTAAA AACTCAGCAG TAGCCATGGG AGAAGTCATC CTGGCTGTCT GCCAGCGCGA TTGCATCACA ACCATCAAAC ACTGGATCAC CATCATCCGA GCTCGCTTCG AGGAGGTCCT GACATGGGCT CTGGAAGAAC TTCTGGCATG GATCCAGTGG GCTGAGACCA CCCTCATTCA GCGGGATCAG ATGGAGGAGA TGACTGGCAA ACAGCCTGAC GTGGACGGGG TCACCAAGAC ATACAAAAGG AAAAACATAG AGCCTACTCA CGCGCCTTTC ATAGAGAAAT CCGGCAGGGG AGGCAGGAAA ICCCTAAGTC AGCDAACCC TCCTCCCATG CCAATCCTTT CACAGTCTGA AGCAAAAAC CCACGGATCA ACCAGCTTTC TOCCCGCTGG CAGCAGGTGT GGCTGTTAGC ACTGGAGGGG ATCOGGCAGA CCCTAAAAAG TCTGCCAGTC GCCCTGGGAG TCGGGCTGGG AGTCGAAGCG GGAGTYCGAGC CAGCAGCCGG CGAGGAAGTG ACGCTTCTGA CTTTGACCTC TTAGAGAGGC ATTGCTTGTT CCGACACTTC AGAAAGCAGC GCTGCAGGGG GCCAAGGCAA CTCCAGGAGA GGGCTAAACA AACCTTCCAA AATCCCAACC ATGTCTAAGA AGACCACCAC TGCCTCCCCC AGGACTCCAG GTCCCAAGCG ATAACACTGT CTAAGCACCC CCAAGCCACT ATCCACTTTG AAGCAGCACC AGCAGCGTCT TGAAACGGCC TTGTCAGAAC TGGTGGCTAA TGCTGAGCTC GAGCCAATCC CGCAGAACAT TGACCGAGTT AAAGCCCTTA TCGCTGAGCA TCAGACAITT CAAAGGAAAC TGAATGATGC CTTGGATCGG CTGGAGGAGT TGAAAGAATT TGCCAACTTT GACTITICATO TICTOGAGGAA AAAGTATATG COTTOGATGA ATCACAAAAA GTCTICGAGTG CATCCCAACA AGGATGCGTA TCGACCAACA ACCGATGCAG ATAAAATCGA AGATGAGGTT ACAAGACAAG TGGCTCAGTG CAAATGTGCA AAAAGGTTTC AGGTGGAGCA GATCGGAGAG AATAAATACC GOTTCTTCCT CGGCAATCAG TTTGGGGATT CTCAGCAGTT GCGGCTGGTC COTATTICTICC GCAACCOTGA TGOTTCGCGT TGGTGGAGGA'TGGATGGCCT TGGATGAATT ITTAGTCAAA AATGATCCCT GCCGAGCACG AGGTAGAACT AACATTGAAC TTAGAGAGAA ATTCATECTA CCAGAGGGAG CATCCCAGGG AATGACCCCC TTCCGCTCAC GGGGTCGAAG GICCAAACCA TCTTCCCGGG CAGCTTCCCC TACTCGTTCC AGCTCCAGTG CTAGTCAGAG TAACCACAGC TGTACATCCA TGCCATCTTC TCCAGGCACC CCAGCCAGTG GAACCAAGGT IATCCCATCA TCAGGTAGCA AGTTGAAACG ACCAACACA ACTTTTCATT CTAGTCGGAC ATECETTISET GOTGATACEA GEAATHAGIT CITECECOGSE CICCACAGGI GECAAAACIA ATGGATTTCT TCCGGCGCAT TGATAAGGAC CAGGATGGGA AGATAACACG TCAGGAGTTT ATCGATGGCA TITITAGCATC CAAGITCCCC ACCACCAAGT TAGAGATGAC TGCTGTGGCT GACATTITICG ACCGAGATGG GGATGGTTAC ATTGATTATT ATGAATTIGT GGCTGCTCTT 349 9 8 2 9 5 တ္တ 15 25 55 35 45

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GETGGGCCGC GGCCTCTGGA GCTGGGATTT GGGAGGACAC AGCAGGCAGC GCTGGCCTTC TOCCTCCTTC COGGACAAGC CTOGCCACCC TCGCTGTGAT GACGAGCTGG CTGATTGGCC TOCAGGGATG GOCCAMANGOT TOCGCAROOG COCGTTOCGG GACCTGCCCA GOGTOCTOCC

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(A) LENGTH: 1830 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDMESS: double
(D) TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 129:

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CGITAAAITC TIGATAGAIT TITTATTAIT

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GOGATIGGECE AGGOCEAGEG CATTOTIGEAE TOGTTTACTT TAAAATGTAE AGATTETTET TCACTOTATC TYOCATCAGA GACAAAGGAG GACCCGCTTT AGCCCTGCTG CGGGAAANTGG

1830 1800 1740 1680 1620 1560 1500 1440

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO:

130:

8

(A) LENGTH: 1864 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

ACGGGTTGCT GCCCAAGTGC ATCATGCAGG CCACGGACAT CATGCGGAAC AGGGCCCAAG GCATOCAGAG GAGCACCCTG AGCGTOTYCC TOGAGCAGGC GGCCATSTTG GCACGGAGCC CCTCTACCOC CTCTGCCAGC CGCCGGTOGA TGGGGACCTC TGAACACCCA AATGCCCCAC GGTGGAGATT CTGGCCAAAA ACCTGCGAGT CAAGGACCAG ATGCCCCAGG GTGCTCCGCG Œ SEQUENCE DESCRIPTION: SEQ ID NO: 129 90 240 180 120 420 360 6

CAAT

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GACATEACEA GCEAGAGEAE AGGAAGEEAE CETTGETTGET GGGGAGGAGG GACCEACAEA COGGOOGIGG CTICIOIGAI TIATITICII GAIOGIAACI ICICAGAGCA GOOCRATIGG GOGGATOGAG CCCCACCTGA GGTGCCGTGT CACACGGGTT AGAGGGTCAC TGGGAAAACAC CICICITICA GIGACIOGCI ACAAGGGCCI GAGAGGTGGC CAGCCAGGGT TOGAGCTGGA TEGACIOTOA TICCITECCE CACCETECAT TETECAGEGG TIGGEEGGIG TIAGAACIEG AAAAANGIC ATIGITICCI GITIGITAAN ATIAGGGIIG TAAGGIGICG TITIGAGGIA TGTTCCTGCT TCTACTAAAA AAAAAAGAGC ACAAAAGAAA AACTAAATTA TTGAAAAAATT

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1140 1080 1020

AGECCECTEG GEAGTTTGTE CECCEAGETT COGTATGCET TEAGGGAAAAG GTEACAGETG

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CIOCTROCCO CAGGOCTIGGG ACCITIGAAGO COTOCCGGCA TOTOGCATOC GAGOCTICCOG GOGAGGAAGC GOOOGACGC CTGTCACCCC TGGCAGGTGG TGAGTTCAGG TGGGGGCTCC

CCCTCCAGGG TGCGCTTCCC TCTCTTGCCG CAGCATACAC GAGGGCAGGC AGTGGCCTTG

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ATGATGCTTT TTTCAAAAAA AAAAAAAAA AAAAAAAAAC TCGAGGGGGG GCCCGGTACC agatacaaga tigaatotot atticitaaa aatacaacti tototiotac titgaaataa 1140 1080

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AACATTIGIA TACCIGAACT TATTITAAAG ATGAACIGAA ATGCACATAG CCAAGICTIG

TGTAAAAACA TICAGTYGAG ACCATATGCA TITICIGIGC IGITIGIACI TGAGGIATGI

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OCTOTOTOG TTTAAAAGCA GATTACAGAA ATGTAAGTCA ACTTAAGAAC RGTGAATGAA 960 900

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GITGITGGIG TIGICAAAGI

GAGATTTOCT CIGACTITAT THATATOGCA TGAAATCICT GOTTTATITT GOGATTITTT

TIGITITITIC TANAGITIGIG TGATTATATA TITIGACATTI

960 900 840 780 720 660 600

AGAGECACAE CAGETETIGGA CATEACEGEE CETIGGAAETIG GGGCCAECAG CEETIGGGCAE TOUCTOTOTO GOGANAGGAG AGAGGGGGCC TOCATOACGC CTGTTACCAG AGGATOCCCG

TACATTICAA AGAAAGGTAT GTIGICTAAC AGGGGACCAA CAGAAGGTAG TATICACAAC

TCTARTITTA ARACTGTATG AGGACTITGT OCTGARAATA GAGTATTITT TTARAGTAAG 840

8

ACAATTTIGT TAAATTACCA AGITIGGTIT TIAATAATIT CICAATATIA TOOGCCAAGA 780 720

ACAGATITIGA CATTACAGCT AAGGAAATAA TITGAGITIGA TICAGAAATC CIGGCATGIG

CITICCCOIT ACIGCOTTIT CACCACCIOT CITICCCCATO CITTATTIAT CIGIATGAAC

660 600

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CITAGCITICC AGGITICACCC TAACCCIGIA CITAACCIGCT IGGIGGACIT GGAAAAGACI

AACCECEGGTT TECTTECAACG TECAÇATTEE AGGTGACEAE ACGTGTCTEE TECTECTEAT COTCATOGAG GCCACATOGG GCCACCCOGC TCCCTCGGGA TOGCTCCGCT GCACTTITTGA AGCTGATGGG AGCAGCTGGT GCCTGGCCTT CGGCTCCTGC GTCCCCAGAA CCCAAGGGAA

CTGGGCCGGC CCATTCTTCA CACGCCTGCC AGAAGCTGGA GGGGTGCTGG AGACCCATAG

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CTTTTAGAGA AAAGTCTGGA CTCAGCCACA AACTCTAATA AGACCTGTAC ATCTGAGAAC

GACCACTOGO TICTITICAGA TOTTICTITOC AAITAIGATA ACATGAGAIT TOCTOTICTO

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TOCAGOTOCO CTOCOCACCO TOCTOTTOTT TOCAAAAAAA ACCCAAGOAG GOCAGGOTOA

CTTOGTCTAA TICGCACTTT CCTCACGAGA ATTAAATTAA GCAAAAAACA AACAAACATA GIGGOCCCIC GICTAGANCA IGAIGIGCCA GITICIGAGA CAICITITIA AGGCICITAC

> 420 360 300

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GACAGACCTA CAACCCGTGG CGGATATGGA GCCCCACGAT TGGAAGAAGA AGCTCGGACC GCACAGAAAA CICCCCIGCI CCICACGCIC CCICCACCIC CAGICCAGCI GACGACIIGG

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COGCCCACC ITCTCTGATC TGCGGCTCA GCTGCATGTG ACCCCAGGCT CAGCCCAACA

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300 360 420 480 546 900 99 720 780 840 90 960 1020 1080 1140 1200

> ACTRICTORGA CAAGTICCAGG AGTIGGATGGT GOCCTACCTG GAGACGCGGC TGGCTGACTG GATECACACC ACTIGGGGGCT GGTTATCCCA GATCACTGAA GCTGAGATGG CTGATGAAGT

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AATTITOCAGI GAAATTITIAA GCGACTGIGA CICTGCTGCA AGITCCCCAG AICTIGAGGA GCTGGAAGCT ATCAAAGCTC GAGTCAGGGA GATGGAGGAA GAAGCTGAGA AGCTAAAGGA

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GOTGATICATO TECATTGAGO AGAAGATGGA GOCTGATGCC COTTECATET ATOTTGGCAA

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GCTACAGAAC GAGGTAGAGA AGCAGATGAA TATGAGTCCA CCTCCAGGCA ATGCTGGCCC

CAACCOTOTT ACCATACTOT OTGACAAATT TAGTGGCCAT CCCAAAGGGT TTGCGTATAT AGAGTICTCA GACAAAGAGT CAGTGAGGAC TTCCTTTGGCC TTAGATGAGT CCCTAITTAG AGGAAGGCAA ATCAAGGTGA TCCCAAAAGG AACCAACAGA CCAGGCATCA GCACAAGAGA COGGGGTTTT CCACGAGCCC GCTACCGCGC CCGGACCACC AACTACAACA GCTCCCGCTC TODATICIAC AGRECITITA ACAGCAGGCC COGGGGTCGC GTCTACAGGG GCCGGGCTAG ACCOACATCA TOOTATTCCC CITACTAAAA AAAGTGTGTA TTAGGAGGAG AGAGGGAAA MCCTTGATGG AAAAAAATA TITITTAAAA AAAAGATATA CTGTGGAAGG GGGGAGAATC CCATAACTAA CTOCTGAGGA GGGACCTGCT TTGGGGAGTA GGGGAAGGCC CAGGGARTGG GOCAGOGOG TOCTTATTCA CTCTGGGGAT TCGCCATGGA CACGTCTCAA CTGCGCAACT OCTROCCCAT OFFICCETOC CCCACCCCAC CCCTCTTCTC COGCTCCCTO CCCTCCAGA PROCEIGNG ALCHAITHTG TITICCTITING TOTITICITITY TOTICITITICA GIGICITICS TTCCAGGITT CIGIAGCCGG AGAICTCCG ITCCGCTCCC AGGGCTCCA GIGIAAAITC CCCTTCCCCC TGGGGAAATG CACTACCTTG ITTTGGGGGGG ITTAGGGGGTG ITTTTGTTTT

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TOTGGACTAT GGTGCAACAG CAGAAGACT GGAAGCTCAC 1TTCATGGCT GTGGTTCAGT

AGCCTTCTTT GTCTTTGGGG CTGCACTGTG TGCTGAGAGT GTCAACAAGG AGATGGAACC

ACCETTCACE CAGGIETICOS ATGAACTITIT TCAAGGGGGC CCCAACTGGG GCCGCCTTGT

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	tcatttccag gggtgggaat tittttitaa tatgtgtcat gaataaagtt gtttttgaaa	1800
	akuadadaa adadadada adadadada adadadada adadadada adadadada	1860
5	эргэ	1864
10	(2) INFORMATION FOR SEQ ID NO: 131:	
15	(i) SEQUENCE CHARACTERISTICS: (A) LENTH: 2041 base pairs (B) TYPE: nucleic acid (C) STRANDENESS: double (D) TOPOLOGY: linear	-
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 131:	
20	GOCACGAGO COCOGCAGOS CCCTGGACCC GOGCGGCTCC CGGGGATGGT GAGCAAGGCG	9
	CTGCTGCGCC TCGTGTCTGC CGTCAACCGC AGGAGATGA AGCTGCTGCT GGGCATGGCC	120
<b>V</b> C	INSCHOOCCT ACSTCGCCTC TOTITIGGGGC AACTICGTIA AIATGAGGTC TAITCAGGAA	180
3	ANTIOCTICANC TAAAAATTICA AACCAAGATT GAAGAGATOG TTGAACCACT AAGAGAAA	_340
	ATCAGAGATT TAGAAAAAG CTTTACCCAG AAATACCCAC CAGTAAAGTT TTTATCAGAA	_8_
30	AAGGATCGGA AAAGAATTTT GATAACAGGA GGCGCAGGGT TCGTGGGCTC CCATCTAACT	360
	GACANACTEA TEATGGACGS CEACGAGGTG ACCGTGGTGS ACAATTTETT CACGGGCAGG	_420
35	ANGAGAAACG TGGAGCACTG GATCGGACAT GAGAACTTCG AGTTGATTAA CCACGAGGTG	480
G.	TOGAGCCCCT CTACATCGAG GTTGACCAGA TATACCATCT GGCATCTCCA GCCTCCCCTC	-540
	CAAACTACAT GTATAATCCT ATCAAGACAT TAAAGACCAA TACGATTGGG ACATTAAACA	00
40	TOTTOGGGCT GGCAAAAGGA GTCGGTGCCC GTCTGCTCCT GGCCTCCACA TCGGAGGTGT	
	ATGGAGATCC TGAAGTCCAC CCTCAAAGTG AGGATTACTG GGGCCACGTG AATCCAATAG	720
¥	GACCTCGGGC CTGCTACGAT GAAGGCAAAC GTGTTGCAGA GACCATGTGC TATGCCTACA	780
}	TRANCCAGEN AGGOSTOGIA GTGCGAGTGG CCAGNATCTT CAACACCTTT GGGCCAGGCA	840
	TGCACATGAA CGATGGGGGA GTAGTCAGCA ACTTCATCCT GCAGGGGCTC CAGGGGGAGC	_00_
20	CACTCACGGT ATACCGATCC GOCTCTCAGA CAAGGGCCTT CCAGTACGTC AGCGATCTAG	
	TEANTGCCT COTGCCTCTC ATGACAGCA ACGTCAGCAG CCCGGTCAAC CTGGGGAACC	1020
\$\$	CAGAAGAACA CACAATCCTA GAATTTGCTC AGTTAATTAA AAACCTTGTT GGTAGGGGAA	1080
}	GIGANATICA GITICICICO GAAGOCCAGG AIGACCCACA GAAAAGAAAA CCAGACAICA	1140
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GITTAAAGAA AGACITTAAC AGGIGICAIG AAGAACAAAC IGGAAITITCA TICIGAAGCI

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AACCAAAGCC TOCCAGAATA AAGAAAGGAC GGACTCGCCA CAGCTGAACT CCTCACTTTT

ACTACCATTG TACACTIGAT GGGATGTATT TITIGGCTTTT TITIGTTGTC

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3 8 35 မ 25 20 5 5 8 8 S CACITITITIGA ATCTICITITI TIATOTAAAA TAGCOTAGAT GCATCTICIGC GTATTITICAA GCTTTATGIT ICICTITINA TICAGAGITT TICCAAGGIC TACTITICAG TIGCAAACIT TOCTITAATG AAATGGATGT GCCTAAAAGC TCCCCTCAAA AAACTGCAGA TTTTGCCTTG CTOCGTATCT GGGGCGGGG CAGGTTGGGG GGCACAAAGT TAACATATTC TTOGTTAACC GACTITGAAA TATICCIGIT GGICAIGAIC AAGGATATIT GAAAICACIA CIGIGITITIG TIGICAGGIG GIGGIGGGC GGCAITGAIT ITAGGGCAGA TAAAAGAATI CIGIGIGAGA AAAAGTOOGT ACTTAATAAA TGAGTOGTTA TACTATOCAT AAAGAAAAAT CCTAGCAGTA GITICITIGI GAAGCIGAAA AGGAACATIA AGCGGGACAA AAAAIGCCGA TIITAITIAT GTTTTTTAT CTTOCTOTGA GAGCATATOT TOTGACTOTC GTTGACAGTT TTATTTACTG GACCOGAGCT GOGAACGGGA ATGGCACAAC CAGTCTGAGA AGGACAGTTA TGGTTACATT TACCAAGCTG CAAGAATCTA CTATATCATG GCAGAAGAAG TAGAGTGGGA CTATTGCCCT 2 CCACTOOCTO CTGAGCCTGG TGAGGTGGTC ACTTATCAGT GGAACATCCC AGAGAGGTCT ATCTTOGGTC CACTTATCAA AGGTGAAGTT GGTGATATCC TGACTGTGGT ATTCAAGAAT TACACTGATG GTACATTCAG GNTCCCTCGG CCAAGGACTG GACCAGAAGA ACACTTGGGA TICCIGAGCA ACAAGGAIGG GCICCIGOGI ICCAGAIAACA AGAAAGCIGI AIICAGGGAA GOCCCIGGOC CAAIGACICI GCIIGIGITII CCIGGAICIA IIAIICIGCA GIGGAICCCA AATGCCAGCC GCCCCTACTC TGTGCATGCT CATGGAGTGC TAGAATCTAC TACTGTCTGG TCAAGGACAT GTATAGTGGC CTGGTGGGGC CCTTGGCTAT CTGCCAAAAG GGCATCCTGG INFORMATION FOR SEQ ID NO: 132: Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 132: 909 (A) LENGTH: 2012 base pairs TOPOLOGY: linear TYPE: nucleic acid
STRANDEDNESS: double 2040 1740 1920 1860 1800 1620 2041 1560 240 540 480 420 360 300 180 120 60

> 4  $\frac{3}{2}$ မ 25 8 50 3 2 5 GAAGAAGOCA ATOTGAAGAT GCTGGGCATG CAGATCCCCA TAAAGAATGT TGAGATGCTG TTCCTCTATC GGAATGGCGA GAACTACCGG GCAGATGTGG TGGATCTGTT CCCAGGGACT GOGAAACTOT ATGCCAACCT TAGGGGTOTT ACCATGTACC AAGGAGAACG AGTGGCCTGG GOCAGTATTA ACCTACAGGA TGAAACTTTC TTGGAGAGCA ATAAAATGCA TGCAATCAAT GATGAAAATA AGTETTGGTA TTTGGAGGAA AATGTGGCAA CECATGGGTC CCAGGATECA NAGCCCCATG GAGGACGGAN TGACATGGAT CGGGAATTTIG CATTGTTGTT CITGATTTTT TATATTTCCT TCTGACACTT GGAAGGTATT GAAATTTCTA GAAATGTATC CTTCTCACAA GOGGCATGGG TGGTGGAGAA GCAGAAGGAG CAATCAAGCT TATCTGGATA TTTCTTTCTT GACAGCTTCA AGCTTCTGTC TITICAAACAG TAACATCTGG AGCCTGGAGA TATCCTCAGG GCCICIBITY IGGITGCCAT TAGIGICACC CITCIGCICG TIGITCIGGC ICTIGGIGGA TTAAGCCCTC TCACCGTCAT CACCAAAGAG ACTGAAAAAG CAGTGCCCCC CAGAGACATT ACTGACCATG TCCATGCTGG CATGGAGACC CTCTTCACTG TTTTTTCTCG AACAGAACAC TITIGAGOTTO TOGAGATIGOT GOCCAGCAAC CCTGGGACAT GOCTGATGCA CTGCCATGTG TACATOCTOG CCATOGGCCA AGATOTOGAT CTACACACCA TCCACTTTCA TOCAGAGAGO TAAGAATATA GOCTTGATGG GAAATTGAAG GTAGGCTGAG TATTGGGAAT CCAAATTGAA ATTICACTIT GAACIGAGGC CAAGIGAGCT GITAAGATAA CCCACACTTA AACTAAAGGC TATTTATTTT ACATOGAAAT AATATGATTT CACTTTTTCT TTAGTTTCTT TOCTCTACOT AAGCACATOT GTAGTGCACT COCAGCAGGC CATGGACTAG TCACTAACCC CACACTCAAA GTOGTTTGGT ACCAACATCG ACAGAGAAAG CTACGACGCA ATAGGAGGTC CATCCTGGAT AACCTOTOGO ACTGAAAGGA ATOTTGAGTT ACCTCTTCAT GTTTTAGACA GCAAACCCTA TGAGCATOTA CAACCTCTOG AGCTAGAAGC TCCTCAGGAA AGCCAGTTCT CCAAGTTCTT GGGCACCTOG CACTAAGGGA GTACCTTATT ATCCTACATC GCAAATTTCA ACAGCTACAT TCCATTAAAG TACTTGTTAG AACACTGAAA AA TITICATICE CCTTGGCAGT GAACTACTTT GAAGAAGTGG TCAATGGGTT GTTGCTGCCA ANGINGAMAA CICATIGATI GOGTITICTAC TICTITICANG GACTCAGGAA 1980 1740 1680 1620 1560 1500 1440 1380 1320 1260 1200 1140 1080 1920 1860 1800 1020 2012 600 900 840 780 720 660

Ξ SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 133:

(A) LENGTH: 1669 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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PCT/US98/04493
WO 98/39448

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8 120 180 240 8 360 420 480 240 8 660 720 780 840 8 960 1020 0801 149 1200 1260 1320 1380 1440 1500 1560 1620 1669 GAGCAGTATT TTAACCAACT TGTATTACAG ATGTTACAGT TCATGTTAGG AAGTCAGAAA AGACTITIGIT TGTCTTTGTT CTGCTGATGT GAGTCATGTT TTGTGGGGGTC TTCCATGGCA CATTIACCTG TTGCTCCGTC CAGATGTTGA GGGCCAGTCT AGGCTGACAC ATCCTACCCG AGGACAAGCC TGTTCTCCAT TTCTTCACTC TCCCTCCCC ATATAGCAAC TCTCCCAGGT MAGATMACC GITTICGACG ACAGAIMAAC CAAAAAIGCC CCACACAGGI ITTAIMACIG FINIATACTA TACTITITAAC AGTACAGACC CTAAAITITTA TIATITIGITIG CTCCCCCAAT CIGATACCAA ATGITTAAAG TIGITIGAAA TCCAAACATG GTAGTGITCA 1GGGTAAATA ITITICIAGGC TATGTAAGAG ITAGCAGCCC ATAGCATAGA AGTAATCAAG TAGCATCTGA GACTIOTTIGGA GCCACTAGGG CCTCTCTGGG CCTAACAGCC TCACTTCCCC AGCCTCACCT CCANGIGIGG ATGCATGAAC ATGAGITCAT TITATGITIT ATATAGCITIT CITAGACATA CCAAACCATC TCATGAAGTC AGTITAATTC GAAACAAAA TGAGGGTGTT TTCTCTCCTG TTAATCTAGA GTGTGCAGTT ACACATGTGT GGATAATTIC ATGITCCAGG GGCGCTTGGC ATCTCCCATG GACTGAITCC CAGGAAGAAA AGCCCAAAGG GAAACCCACG ATTCCTTTCG AGTAGATGTG GGAAAGAGCC CATTGGAGGA TATCAGGTCC TGTGAAATTC AGTTGTGTGT GTGGCTCCTT GTTAGCAGTC ATGTTGACAT TAACCTTGGG ACACAGTGGG TTAGCCTGGA GAAAATGAGA GGCCCTGCCT GGACCCAGGG AGAGGAGCCA GTGACACAGG CAGAGCGGTG CAGCCCTCCT TCCCTTCCAT TTGGAGGAGG CATAAAGITC TATCAAGGAG TTCTATCAAG GCATCCATGT CAGTGGTGCT ATGCTGGTTA SCICITIAGGA GOCICCCCAT CCACCCITIA CATGATISTAG GGACCAGIGI CITISTGAGAI IGOTOCCAGO AGCCTGCCCG CTTACCTCTG CTGAAGCATA AGTGGACTTT GCTTTTGGGG CITATICICIO ATACATICCIO GAGCICIOCC ICTICLACIGIC TAGATIGGAC CIGGAATICIC ICATCTACCT CTTAGTCTGT CAGTTTCTAC GTGTGAGAAG CAAGCTTGTG GGCCAGTGTC CTIGIACATG CIGIAGCACT TAAAAAIAA TICCAGGGTT CCCTGGAAAA CCAGTCCCAG GGTTCCTATG ATCTGTAGTT TCTACCTGGA TTATAACTGG TTTTGGGTAC CTGAATTTTG ATTGGTTAGC CTTAATTATA GTCTGGCGTG ATCATGTAGA ATCTTTTCTG GTGAACAGAT DECIGICATE TORCACTOCE ATCAGGGCTO TTAGTGGCAE CTGTATGAGG ATTCATAAAT CAGATAAATT ATTCAGTTTT TGTGTTTAGA AAGCTAAGTA CAACTIICAGA TITITICAAAT AAAAAITIIG TCATAAAAAA AAAAAAAA (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 133: STCCAGGGA ACAGCACAGG TTAATGCGTC TCCCTAGAAC S 2 15 ຊ 25 8 35 <del>수</del> ₹ 8 55

CACTITITISCT ATATAACCTA AGICATAACC CICTITITAGI TACCIGCCAA ACTCIGGACT TICCTITIGEA CACTITICATO GAATTCATCT GITAACCCCI CITCCITIGA GAGAGCACCG GCGATGGIGG ITAACICCIT GIGITITICIC ICTOTOCTAC TOGITATICT TOAATTAAGO ACAGACTOOT CAGCTOGGTI GCTTTAICAT TACCTTCACT GACCAAAAAT TAAGGAAGGA AAACACAGTT TTTAAAAGGA TCCATCTTTT AACAGCCGAA COTOTOTOAA CGATCATITIC TGACTIAACC GTGAGATGCT CACGAGTACC CTTCCTGTTG TTTTGTTAGC ATTGAAATCG AGACTATTTA TTTGGAATAT ATACAACAGT GTTTTTCCAC TGTATTTCAT INCCADAGOT IGAGAACIGC ITICICIACC ITITIOCADAA IAAITGAIAT ICCAIAITIGG ATTOTICAAAG ACTTOGATAT GGTGAACCTA TTAAACCTAG AAATTGTATT CATCOTTTCA TCACTIGTIGG CTGAGTTCCC CAGCCCCTCT CCTCCTTTTT TTTAGATGAG ATTTAGCACA CICICAGITA TITAAACATG CAACATITICT TGAGTATGTA TGITGAGGCC ATCIGAGCTC ATAGCTGATT CAGTAACCAG ITTCATGCTG TGTCATTCAC ACTCACTACT TAATACTGCC ATGGTGAAAA TGTGGAGGAA AAATGTATCC ATGTGTGTCT GGGAAGCATA TACACTTGTA CATTITITAA TACTCIGAIT CIGIAACAIT TCIGAGIITIT GITITIGITIT ACAGAAAAA AAAAAAAAGT GATAAAGCAA TCAGAAGACC AAGAGGTTTA CTATTGATGC TTAGGGTCGT CTGACCTTGG CTGGCCAATA GACCTACACG GCCAAATTAA TTTACGAGAG TAATAATTTT ICAAAAGCCA ATITITITIC TGTATITICT GTATGAAACT GCCAATATCA TGAATAGAAA GGGAGAACCA TAAAGGAGAA AGAACGIGAT GTICTGTIAT GTICATGIAA ACCTAAAGAA ACAGTGTGGA GGCAGGCGG ATCAGCCGAA CTCTAGGGAC TTGGTGTTGC TTGGAAGGCA ICCATACCTG CATTITISCAT ICTICGIATG TAATCATATT GCCAAAGACA AACTATITICA GIGICTITIT CCTGTTTCTG AGGATTTACA CATGGCATTC AGTGTTCTGT ATAGATCTGC CTACCTTTGT GAATAATOTG TGTGACCTTG CAGTTCTTCC ACAGTTCAGC AAACAAGTGC ACCGATGTOT CTATGGTGCT GCACCTTGCT GTTGTACTTC TGAAATCAGA SEQUENCE DESCRIPTION: SEQ ID NO: 134: TGGTTTATAT TGCAGTTAAC ACAGTTACAA AGCTGTAATG ACCGRATCTC TAAATCAAAG TATATACATT GTGTGGTGTT LENGTH: 1565 base pairs STRANDEDNESS: double TYPE: nucleic acid SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 134: TOPOLOGY: linear 3 8 0 8 Ī £ 2 2 2 ឧ 25 35 ಜ 5 45 င္တ 55

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720

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PCT/US98/04493

TCATTIATIG TAMATAMCAC TITTICCCCMG ACCTACCATA AAGTITICTGT GATGTATTGT 

1560

1500

CCATOTOCTC CCGTGCACTC CTCAGATGGT CAGAGGGGTA ACCCAAGTCC TTAGAGAATT

1320

TOGGGACCAA TAGAATATGT GATGTGTGAA TTTTCTTTAA AAAACTTAAG GAGTCTTTGC

1565

(2) INFORMATION FOR SEQ ID NO: 135:

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CTCCA

Ê SEQUENCE CHARACTERISTICS: LENGTH: 2007 base pairs TYPE: nucleic acid

**60 €** € TOPOLOGY: linear STRANDEDNESS: double

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TOTAMAGEC COOTTATACE COACTITIOTS CAGCAMAGAT COCCOTOCAG STOACAGCOT

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SEQUENCE DESCRIPTION: SEQ ID NO: 135:

OCTOTOTTOG TOTTOGGACC AGCAGAAGGC AAACGTCCAG CCAACACACA GGACTGTAAG CATTITOTOGO CAGGOTOGAC AAAITCOTGA GGCACAACTT GGCTTCAGIT CAGATTICAA

180

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AATTATTTTC TCTGTATGAT TAAAAGT

GTCGCCAAGA CATCTACATT GTAAGAGAAC ACAGTGGAAG ATCCTGTCCT GATTCTCAAA TATANACTOT ATANAAGGIT CIGITITITAA AGGIGGATITI TCATICCICI GGGGACAGIG TITITITITA AATITAGGAT AACACATTIT TGITICTAAA GIGATTIGIG ATTIGIGCIG

1980 2007

1920 1860 1800 1740 1680 1620 1560 1500 1440

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CHOTAGCITT TIMAMAGGAA ACCUMGICAT CCCACTATGA ATCTGGCATC TICTIATGCT TIGACAGGIT GOGCIGIGIG TOTOCOCATO TOTOTATACA TITICCAGGCG TOCCIGIGIC TACCTICTOC TIGTIGAGIT GITTIGGCAT ICATATIAAA AGCCAGCAIC TCACTATITA

TTCAGGAGGG GCTGGATCAA ATTTTGAGAG GCTTATGGGA AAGGGAGGGG GAGAAGAAAT TCTAGTGTTT TGGCCATACA TCAACCAAGG GGTTTAATTT ATCCAATGCT TGACGACATG

GARTAGAAAC TGATAGCATT AAAATACTCC GTTCCTCTCT CTCTTCTCGC TTCCTTTTTT TGACATTTAT TITATTATTT ATTITAAATG TITACATCTT CTITATGTTG TATCAAGCCT

120

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30 GAGCATATICE TGAGTTTTIAC TICCTTATIGG CTTGCCCTCC AAGTTCTCTC TCTCATACAC CTTCCCCAMA GTCAGAGGTG ATTTGATTTG GGGAAGACTG AATATTCACA CCTAAGTCGT

> 360 300

> > 30

(2) INFORMATION FOR SEQ ID NO: 136:

(i) SEQUENCE CHARACTERISTICS:

(A) LEWOTH: 1291 base pairs
(B) TYPE: nucleic acid
(C) STRANDEINESS: double
(D) TOPOLOGY: linear

AGGACTOTGA GOTACGTGCC CTGTGAAGAC CCCCAGGCTT TGTCATAGGA GGTCGTTCAG

ACACACACCC TIGCICCAGA ATCACCAGAC ACCITCCATGG CTCCAGCTAT GGGAACAGCT TOCCTITICTS TITTGGCTTAG GAACTICTGT OCTICITOTO OCTCCACTOS 480 420

3GGGCTGCAGG CAGCTCTGGG ACOGCACAGG

TGATCAGCTC GTGTAAAACA CACCGTCTTC TTGGCCTCCT GGCAGTTCTT

8 TOTOCOAATA GTOCTOTOCO TOGCCAGTTO AATOGGGGAA GCTOCTOGCA CAGGAAGGAG OGCTGAGGCT TAGGAAATTG CTGGAGCCGG CTCCAAGCAG ATAATTCACT

> 720 660 600 540

780

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TITICICATIT TIMIGCIGIT GOSTCTIAGT TITIAAATIG ATATAAAGAA CICAGCAATG

240

180 120 CTTTTNACCC TCCCCCTTCA CACACATACA TATCAGGITG TITTCTAGTT AAAAACCCAA

GIAGCICAGA TICTACTITA AIGICAGIGC AGATITICCAT IGAAICAIGC CATTAIGITI

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SEQUENCE DESCRIPTION: SEQ ID NO: 136

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4 AGCATOTOM AGTOMAMOS CATOTOGGO TGOTGOTTOT CITTOTOMG CTOTGGGGM TCAGAGTCAA ACATCATTCT GCCTGTKTTO GGGGCCAGGT GTGTCACACA

50 ACTITITITIC TOTTITICCIT GATGGACCAA CAGTGCAAAT GCAATCTCGC CATTTAACTT AGGAATCTCC CTCTCCTCTC ACTTGATTCC AAGTGTGGTT GAATTGTCTG GAGCACTGGG

TCAGGTCGAT TTCCTTTCCT GATCAGACAT CTTTGTGCCC CCTTTAGGAA GGAAAAGAAT

ACACCTACGA TGTGCCAGGC ACTGTGTTAG GCGCTTTTAT ATAGATCCTC GTTAGGATGA

GACTAAGGGA TGAGGACATC TCTTTATAAA AGGCCCCTAA GTAATGGATA AACAGAAACA

900 840

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TOTTCTCAAG GACTTATCCC CTACAATATT CTCCCACTCC ATACTTCTCC TTCTACCCC

CTTAGAGOTG AGAAGGTCTG TCTTCAAGAT CCAAGGTAAG ATTOCCTTCA GTCTGATGTT

1260 1200 1140 1080 1020

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ACATTEMAAT TEGAGMAGGE GACMACEGGC TOTTETCCAG TOTECCTECA TGECAGETETE TCATACAGIT ATTOCCATGA AAGGCAGAAT GITTGITICA AAATTAATCI AGITTICIGI ATAGTAGTIG TITICCAGAAA ACACTICCTC ACAATIGTAC TICCCAATCA AATCATGIGA AACTICAATG GICTACIGAA ACAAAAATGG TAACTITITCA TTAGIOGATT ATTTAGAGTT GITTIATITI CTACICATAC TIAGGGITIA GGAAACACIA CCACTAGITA TCATITAAIC

480 420 360 300

600

CTGATAGACC ACTATTOGCA AACAGTATCT GTCAACTACC AAATGTGTAA AATTTTCTGT

ATTTCACTIT GICTTATITG TAANIAGIGA ACTAAAACIT TIGGCAGAIC AGCAACATIT

360

WO 98/39448

- PCT/US98/04493

-	WO 98/39448	PCT/US98/04493 W	WO 98/39448 P
	361		362
	GCTGAGCCTG TITTITIAAGC TAATGIGTAT TCTTACTAAT GTTCCTATCA AGAITGGATT	099	THETAAGTIA GITTGAETTA CANTANTATT CHCANTINAA ACTAALAATA AAGGETTECAT
	TGIAATATAT GCTGTCTATT TCTAATGTTC ACATTCATAT TTTGAGGTTC TATCTTATTT	720	TTICABAGAT ABATTICABAT TICTICATICOT GABATBACA CCABARTACT GABATTICATIC
5	TAATAGAGAA CAGACTTCTC AAAAAATCTT CAGAAGCAGC TTATTATTGA AATATCGAAA	780	TACATACAG TITICTACAS AGACANCOS AGAATITACA ATTICGAGA TITICAGA TITAATAACTA
	TATTGADATA AACCCGGTGG GTTAGATTAC TCATCTGTCC ACCAAGTGGG ACATTTGCAT	940	CONTRACTA CARABACTICA PHYLAMARIA STOCIALINA SACTARACTA STOCIAL
2	GGACTGGGGG CTTAAAGGAC TTAGAAGAGA CCTGTAAGTA AATOCTGAAA ATGAGCCAAT	01	GITITIC CITATION CALENTINA CONTINUES OF STREET
•	CCCCACTIGA AIGGITACTG GAGIAAACCC ACCITTACCA CCCCAAITAC AGCACCGGAG	096	AACTTACCA APAGAATTTT PACETTAAAA CAACATACA CACETTTTTAA
	GCCGATAAAC CAACTIGGCT CTGGTTCATT TITCTTTTCT TCATITIGTGA TGCTCAGAIT	1020	TOTICATIONS ATTITUDADE CONNECTOR OFFICE ATTITUDADE TEXTORERY
15	CAANATOTOT GITCIACACT GITACAGGCT TCTCITITIOT ITGAITAAAG AITITIAGICC	1080 15	THIGHTCAC AGTIVATOR CONCOCCAA GITHCHATT CAARLAAC GONGAARA
	TACTITITICIA TOGACACATI AGAITATICA GAGACCAAAA TAGAAGAAIT TOCTICITAGA	1140	CATTICTAGT AGTETIANTS ATTITITETS SAARRAGITT STGATTECAA TOCASSIGN
20	TATTITICAG AAGTCAGCAG ATTIGIGGCA AATCATTIAT TIGCCTTITT AAAAATICAT	1200	TICATTACCA TTACCICTAC ACTECAGAAG AACCAAAACT CCTTTATTAG AATTACTTCA
ì	TTAAGGAGIT CAGAGAGIAG ACTACTCAGA AAATTATTTC AGGIAATTGT CTAAGAGGTC	1260	CHICTGIATO 030AAAMA TICTGAAAGG CTAGAATGAT ACAAGTGAGC AAAAGTTGGT
	aatattitit aatgcatait gaatcaaata a	1291	CACTICOCT ATGRACITOR GOTAAGATO TOTAAACATT COAAAAGAC ATGAGTTGAA
25		25	CCTAAACTCC CTTOGGAATC TGGAACAAAG GAATATGAAA ATTOCCATTT GAAAACTGAC
	(2) INFORMATION FOR 6EQ ID NO: 137:		CAGCTAATCT GCACCTCAGA GATAGATCAG CCAGTGGCCC ANAGCCATTT CAAGTACAGA
. 30	(i) SEQUENCE CHARACTERISTICS:	30	ANTHAINGAG ACTACAGCTA AATAAATTIG AACAITAAAT AIAAITITAC CACTITITIGE
			CTITIATIAAGC ATATTITGTAA ACTCAGAACT GAGCAGAAGT GACTITIACTT TCTCCAAGTIT
ž	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	35	GATACTEMOT TEACTOTTCC CITATECCTC ACCCITCCCC ITCCCTITCC TAAGACAATA
ć	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 137;		GIGCACAACT TAGGTTATTT TIGCTTCCGA ATTIGAAIGA AAAACTIAAT GCCAIGGAIT
	OGCACGAGGA CCTACTTITG TAACAGACCA TOGITGIGTC CAAGGTAAAA CCACAGTGAT	09	TITITCITIT GCAAGACACC TOTITAICAT CITOTITAAA TOTAAANGIC CCCTTAIGCT
40	ATTITIOGAT GCTITOTOTO CAALCTIGAC TIGITITIOC AGEATCATIA TICAGACTIC	120 40	TTIGAARTAA AITTICCTTTT GTAAAAAAA AAAAAAAA AAAAAA
	AAATTGTGAA TCTTTTAAAC ATCTTGATAA TTTGTTGTTG AGAGCTGTTC ATTCTAAAAT	180	
45	GTAATGAAAT TCAGTCTAGT TCTGCTGATA AAGATCATCA GTTTTGAAAG GTTACTGATT	240 45	(2) INFORMATION FOR SEC ID NO: 138;
?	TICCTCTICC CICITAGITI TITACCCAAT ATAIGGAGAA GAGTAATGGT CAAICTTAAC	300	(i) SEQUENCE CHARACTERISTICS:
	AITITICITIT AATICITIAA TAAAGCIGCT GOGCAGIGGI GCAGCATICC TACCTAGIGI	360	(A) LENGTH: 1935 base pairs
20	CATAAAAGCA AAATACTTAC ATAGCTTTCT TAAAATATAG GAATGACATT ACATTTTTAG	420 50	(C) STRANDEDRESS: double (D) TOPOLOGY: innar
	GAGAAAGTAA GTTGCTTTGC ACGGCCTACT TAATTCCTTT CCATATATTG TGATACAAAC	480	(At) SEQUENCE DESCRIPTION: OR OF 118:
55	titigaatat ggaatcttac tattigaata gaaatgigta tgtataatat acatacatac	55	TCTGAACTAA TGCTVACAGA TCCCCCTGAG GGATTCTTGA TGGGCTGAGC AGCTGGCTTGG
	ataagcatat atgigigigt gigigigiat atatatatat atgcatgcig tgaaacttga	009	ACCTAGRACT GACTGACATT CALTGRGATG AGGGCACCTT TCTGGRACAG GALTGRAAGC
	CTACACAACA TAAATCACTT TTTAAATTCC AGGAACGGGT AGTCTGACAC GGTGATTATC	099	TCINIGITIT ATAIRCATT TCATCIGIAC TIGGACCICA CITTAGACAA GAGGAAACTA
99	CTITICAGGC TGAATCCOTT ATTAACTIGT TATTTAGGTT TTACTCCCAG TAGCAAGGGA	720 60	

  . 00 

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25 20 5 25 8 35 ಜ 5 S 8 S 8 TOGATIOTO TITICCCCAGG CAGACGGCCC CICTYTTCCC AGCACTICCC TGCCTCCCCC GGAAGCTTAT TTTCCCCOTGG CCAGGATGCA TTTCTCTGAG TGGAAACAGG TTCTTGCATG GACCIACAGAC ACAGATTICC TOCTOGOGGA GGGAGGAGTC CACOCATCCT GATOCTOCCT NOCCCCCTTO GCACAAGTCA GATGAAGCAC GTTCTGCCGG GGAGGCCCTC AMCTTCCAGA TODOTHOOTO GATTTGAGCG GAAAGACTCC CAAAATOTGC CAAGAATTTC CCROTCCCAG GOGGAGGOGG TOTGGCAACG TGCCCCCCGC AGAGGCCACG CATGTTTGAC CAAAGCCCTC GUAGRAGUTG AGRAGOGTGA ACAAACCCCCG AGGGAGGCCC GOCCCTTGCT CCCGAGTTGG CAGGAATAAT TIGAAATGIG IGAGGIGACI CCCCGGAGGC CIIGGGCIIG GGCAITIGGG TCHATCOTCO ACCUACICAG TOGGGAGAAA COTCTCACCG GACAGOTTOG GAGAATGAGG CICCICAKOT GACCACCCIC CICIGACCSA CGCCCCCICC TIGICIGAAA AAAGGAGCCI TOCCCTTCAT CTCCGTCCCC CTCTTTGAAG CTGTCCCCAT CTCAGTGTCA GACCAGCCTT GACTECCACE TOTACCCTT CICCTCACCA TOTCTCCCCC ACCUTACCTC GCAGGOCAGG GGAAACTAAG GGCAAGCAGG ATACAGGOCG AGGGATGTOG CAGGTGAGGG ATTOTOGTCC GAGGACAGCC TTTTCCCCAG OCCTCARAGC ATTOCTCATC COTOCCAAAC AGOCCTCAGO CCAGCACCCA GTICCTCCTC ACATGOCAGG TGAGCACAGA CTICTAGTIG TACCCCAACC CTTTGCGGCT CTGTACACAT TTTTAAACCT GGCAAAAGAT GAAGAGAATA GOTETAGETE CTCTCCAAAC AGCCATGCCC TCCAAATGCT AGAGACCTGG GCCCTGAACC CTUACACCCA AATCCAGAAT CCCTGGTCTT GAGTCCCCAG AACTTTGCCT CTTGACTGTC AAAAGAATGA TOTCTOGAAG GGCTTAAGGG ACACAGTGGA COAGGGGAGA GTCCTCATCT TCAGCGGTGC TGGGGAACAG ATGGAGGGGG CAGTGGGGAC AGGGCTTGGG CAGACACCAG CTOTAGACAG ATROCCCTCAG AATTGGGGCA TGGGAGGGGG GSTGGGGGAC CCCATGATTC COTTOTOTO CTACCTOCAT COATGGAAAA TTAGTTATTT TOTGATOOTT TOOCCTGOOT AGCCACGGAC TCCAATGCCC AGCTCCTCTC CCCAAAACAA TCCCGACAAT CCCTTATCCC (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 139: TOTOGOGIST TAGTISCCANA CTITGAATAGG GGCTGGGGTG CTGTCTTCCA (A) LENGTH: 1446 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 1440 1320 1260 1200 1140 1080 1020 840 720 660 600 540 480 420 360 300 240 180 120 1380 960 900 780

(2) INFORMATION FOR SEQ ID NO: 139:

36 4

TOCAMAGITA GCTOGATOGC TOMAGGTOMC TTAGGTAMGT TOGGAMGTOC ATOCTTOCCA

GCAGAGCAGC AGTTAAACCC GTGGATTTTG TAGTTAGGAA CCTGGGKTCA AACCCTCTTC

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AGAGTETAAA ACETACAGTG AATCACAATG CATTTACCCC CACTGACTTG GACATAAGTG

TAATGGATAA TCTTATTTAT CTAAACTAAA GCTTCCTGTT TATACACACT CCTGTTATTC

25

TOGGATAAGA TAAATGACCA CAGTACCTTA ATTICTAGGT GGGIGCCIGT GATGGITCAT

TOTAGOTAAG GACATTTTCT YTTTTTCAGC AGCTOTOTAG GTCCAGAGCC TCTGGGAGAG

GAGGGGGGTA GCATGCACCC AGCAGGGGAC TGAACTGGGA AACTCAAGGT TCTTTTTACT

GIGGGGIAGT GAGCIGCCIT ICIGIGAICG GITICCCIAG GGAIGIIGCI GIICCCCICC

1200

1140

89 1020 960 900 840 780 720 660 8

TIGCTATICG CAGCTACATA CAACGTGGCC AACCCCAGTA GGCTGATCCT ATATATGATC

AGTOCTOGTG CTGACTCTCA ATAGCCCCAC CCAAGCTGGC TATAGGTTTA CAGATACATT

AATTAGGCAA CCTAAAATAT TGATGCTGGT GTTGGTGTGA CATAATGCTA TGGCCAGAAC

TOAAACTTAG AGTTATAATT CATGTATTAG GGTTCTCCAG AGGGACAGAA TTAGTAGGAT

GICGCACAAT AGGCCGICIG CAAGCIGGGI TAGAGAGAAG CCAGIAGIGG CICAGCCIGA ATATOTATAT ATGAAAGGGA GGTTATTAGG GAGAACTGGC TCCCACAGTT AGAAGGCGAA

AGGCCAAGAG CCCCTGGCAA CCAACCCACT GGTGCAAGTC CTAGATTCCA AAGGCTGAAG

AACCTOGAGT CTGATOTCCA AGAGCAGGAA GAGTOGAAGA AAGCCAGAAG ACTCAGCAAA

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CTTTCCATTA CATGAGCTGT CTCAAAGCCC TCCAATWAAT TCTCAGTGTA AGYTTCAAA

TECCTGAAGT TGCCCTGGTC TCTGCACCTT CTAAACCTAG

TTCTTAAGAG

1920 1860 1800 1740 1680 1620 1560 1500

1935

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AGTOTICTACC ACCAYAGTOG CCATACCAAA GAGGCTACCG

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GTTCAAAAAC CTCAAAACTG GGGAAGCTGA CAGTGCAGCC AGCCTTCAGT CTGTGGCCA

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1440

1380

1320 1260

AAAACTAGCC AGAAGTCTCT TTTTCAAATT ACTTACAGGT TATTCAATAT AAAATTTTTG

TAAAGOOTET GECAGATAGG AAGATGETAG TTATGGATTT ACAAGGTTGT TAAGOETGTA

AACTTICACI TICIATGICI ACCICAAAGA ATIGITGIGA GGCTIGAGAT AATGCATTIG

CACTAATTGG CTATGICICT GGACAAGITT TITTITITIT TITTITITAA ACCCTITCIG

AAACACTATE TICCCATCIG TITCICAAIG CCIGCTACIT CIIGIAGATA TITCATITCA CHATRIGCATIG AGGTGAACAG GACGTAGTIN AGGCCTTCCT GTAAACAGAA AATCATATCA 540

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GTTTCTGCCT CTTTTTGGAA AAGAAAACAA AGTGCAATTG TTTTTTACTG GAAAGTTACC 480 420

360 300

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GIGCCIAGCI TIGGAAAGIC TAGAAIGGGI CCCIGGIGCY TITTIACITI GAAGAAAICA

CTCAGCTCCT CAGGTCAGCA AGTCTACTTC TCTGCCTATT TIGTATACTC TCTTTAATAT

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Š	WO 98/39448 . PC	CT/US98/04493	WO 98/39448	PĊT	PCT/US98/04493	
	365			366		
v	ттста	1446	(A) LENGTH: 497 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear			
3	(1) TARENDAMPTON END CENT TO NO. 140.		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 141:	NO: 141:		
	OF THE CHARLES FOR THE THE CONTRACT TO THE CON	·	INGGACTAAC TIAAATITCIT TTAITCAICT TITAITTAIT AAAAAATITT AITITCITIGA	NIT AAAAAIIII AIIICIIIGA	09	
01		01	ATTICCIGI AATTICCITA RGCTCTICIA TAAAAIGITA TATICAIGIG AACCATACCI	MA TAITCAIGIG AACCAIACCI	120	
	(B) TYPE: nucleic acid		CATTATCCTT AACATTTACT CTCAAAAGC TITTTTATTTT TATTTTTTG AAGGTAGTTT	ITT TAITITITIC AAGGIAGIIT	180	
1	(D) TOPOLOGY: Linear	•	TICTOTOTOT ACTICTOTABC ATGAITTITGC TITCAAATCA ITGITOTOCC CCCATACAAA	ICA ITIGITICICC CCCATACAAA	240	
15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 140:	C! 09	ATGCCTTTTA TTTTTGAGGA TCGTGGACTT TTTAGTATGG CATGAGTGTG CTAAAAGCCA	ICG CATGAGTGTG CTAAAAGCCA	00	
	TITITITI TITITITI CALCULATOR CALC		GATATETITE CACATICACT GGTGGCTTTG ACACCTAGIT TITAATETEC CATECITACT	FIT TITAMICICC CATCCTIACT	360	
20	TAACCCAAAA ACTITCIGIA GACCIGIICC TITGAAACA GCALCACIIA TIGACAGINA	20	TTANACCTO ACADTOCAGT CCTCAGTCAG GOCCAGGACC GGGCTGAGGC CCTTTGTGGA	ACC GGGCTGAGGC CCTTTGTGGA	420	
	AGACTICAGTA TAAAAGCACC AGCATCCCTA CTTGGGTGAT GGGGATTAAT TTTATAGCAT	180	GAIGCIGCAC CACCAGCAGA AGGCIGAGAC CIGGTIACCT GIACCIGITC ACTIGIAAIA	CT GTACCTGTTC ACTTGTAATA	480	
	TOCATTITICO TAGTOCCACA TOTGAAATTO GATTITIGATO ATCTTAATOT ATAITOTACO	240	AAAAGAATTA TCTAAAA		497	
22	CITATAATAA AAGATCAAAA GATATATCTC CTATGAACAG ATTGGAGATA GGAGATGAAA	300 25			}	
	ACTIVOGANGS ATOTCTITAT TCTAATOTGA GGGTAGGGAA AATOTGGATA ACATTACTGG	360				
ç	COTGARGGAG GCATTOTICT TIAGITICGAG TICTCATTIT TATICTCCAG TACTGACTIC	420	(2) INFORMATION FOR SEQ ID NO: 142:			
ર	TOGOGAAAGC ATACTTTTTC ACTIOCCAGGT ACTIGAATGCA GAGGCTCAGT GAAGTATATA	480	(1) SEQUENCE CHARACTERISTICS:			
	TOTOGGAAGT GCATGCATTT COTTTATTAG CAAACATAGC TOGATTAAGA CAAAGTTGTT	540				
35	COTTTGGAAA COCOTTAAAG CCTTAAGTGA ACAAATCTAG CTAACAGTGA ATGAACTAGG	600 35	(D) TOPOLOGY: linear			
	TAATATAACT TOCATATTIT TAATTICCTT TGGTTAAAGG TCCCCCATAC TICTCTGTTC	099	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 142:	NO: 142:		
9	GARGACHIGA GAAGHAIGAT TACTICAGIG TTAGTITICT TAATITITIT TITCCCCIAL	720	ATCHGGCAGA GGCAAGCTGC CTGCCAACCC CCTCCCTCAA GGAATGGCCT TGCCCAGGAA	AA GGAATGGCCT TGCCCAGGAA	09	
€	TIGICCCTIS TCACTITIGIT GCAAGCTAGA AATCTGIGGG TTATACATAG GGCAGCTCTT	780	TGCCCACCAC ACATACCCTC TTCTTTTTT CTAGTCAAAC TCTTGTTTAT TCCTTGGCTT	AC TOTTGPFFAT TOOTTGGCTF	120	
	TOTGAAAGTG GTTTAITCCA CTGGAGAAAG GGGATTGAAA ATCAGTTAGA ACCAATGTAT	840	GCCTCCCTCT TITCCTCCCC TCTCAACCTT TIACITCTGG TTTCTAITTC ATGGGAITTG	GG TITCTATITIC ATGGGATTIG	180	
45	TICTIGCCCC ACGGARCACT ATTCCTATAA GATAGCTGAA AGAAGCTGCT GTGAGGAGCT	900 45	GOOTTGAAGT TAAACTTACA ACAGTGCCGC CAACACCAAG TCTTGCAGGA AAAAAATACA	AG TCTTGCAGGA AAAAAATACA	240	
	CAGCTCCAAA CACAGGATCA GCACCTTGTA TAGGAATTCC CATGAATTAT GACTTCTCAT	096	aagaaaitta acaaaaaaaa aaaaaaaaa		269	
Ş	TCTOTITIAT CAGAGISCAT ATAIGICCTA CITCAGGAAA AGTAAAACAG TCATTTAGGA	1020				
3	AAGAAAGTCA ATCTGTATCC TAAGCATTTT AATAAAAAGT TAAAACAAAA AATTAAAAGG	1080	(2) TANDODAMMON BAD GEO IN NO. 143.		,	
	GACACTICGAG, GGGGGGCCCG AAACCCAAT	1109	(2) ANY CONTROL FOR SEQ AD INC. 145.			
55			(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1169 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double			
	(2) INFORMATION FOR SEQ ID NO: 141:		(D) TOPOLOGY: linear			
9	(1) SEQUENCE CHARACTERISTICS:	09	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 143:	NO: 143:		

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AAAAGCCAAA CTATAGGATA ACACAGAGCC CTTTTTGAAA ATAAATTGGC ATTGGAGTGT Ě SEQUENCE DESCRIPTION: SEQ ID NO: 144: 69

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GTATTITCAT GAATTTACCA TATATCTITG TITITCTICA ACGAAAAAGT TAATTGAGGC

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(2) INFORMATION FOR SEQ ID NO: 144: Ê SEQUENCE CHARACTERISTICS: (A) LENGTH: 1944 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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TATOGAGCTG GGGTTTAACA CTAMAAACTA GAAATAAACA TCTCAAACAG TAAAAAAAA AAAAAAAAC 1269 1260 1200

8 ႘ GACAATCAAA AACGACAACA AGCTTCTTCC CAGGGTGAGG GGAAACACTT AAGGAATAAA GCACTOTTAG GOTTGOTTAC TOTACAAGGG ACAGTTGCAT TTGTTGAGAC TTTAATGGAG ATTTOTOTOA CAAGTGGGAA AGACTGAAGA AACACATOTO GTOCAGATOT GOTGGCAGAG 1140 1080

AAGAGCGGAA ATCCAAGAAC TCCTAGTACT CTGACCAGCC AGGGCCAGGG CAGAGAAGCA 1020

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GCTGCTGTGG GGAAGAAGAA GAAAGATGTG AFCCTGGCTG ACTTACTGCC TICCTTGGCT

STTATOGACA CCACCACAGC CCAGGGCCGA AGCCCTGTGG AGGTGGCCCA GGATGTTCTT GCCTACATCC ACACCAACCT CTCTGTAAAT GCCATCACCG CGGATGGATC TAGGTATGGA

GTTTATCTTC GAACICIGGC TCCTGGGCTC TTCTTCAGCC TCATGCCTCC AGGGCCAGAA

960 900 840 780 720 660 600 540 480 420 360 300

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GICCIGAACC IGGICCIGIG GGCCATIGAA AAGITAGAIC IGIGAICICI GGGGIITITIG CAGACCAGCT TTTGCACAAT GAAGCGCAAG GGAACAAGTG GTTTGCCTGG TGTCCTACCT 25

CCCCCTTGCC TGGTTCCAGC TGTCAGAGGG ATACCATCCT AGGGTCTGGG AATCCAAGGC

CAAGTTATTT TCATGTTACC TGGAGAGTGT CCAGAGGCTG CTCTGAGGCT GAGGTGTGTT

CACGAGACTC CTTOSTITGT GGTCCGAGAT CCTGTACTAA GGAGGGTCTG GCCAGAGGAA

960 900 840 780 20

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CTIGCCCTCCA TGATCAAGAG GAGGCAAGGC CACATTGTCG CCATCAGCAG CATCCAGGGC

TITIGACTOTO TOCOTOCOGA GATOGAACAG TATGAAATTG AGGTGAOCOT CATCAGCOCO AAGATGAGCA TICCITITICG AICAGCATAT GCAGCCICCA AGCACGCAAC CCAGGCITIC GTOGACAAGA GOGTCATOGA GACAAACTAC TTTGOCCCAG TTGCTCTAAC GAAAGCACTC

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ATACTTOTCA ACAATGCTGG

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GCTTCTCATG CCACCAAGGT GCAGACACAC AAGCCTTACT TGGTGACCTT CGACCTCACA AMACTIGGTICC TICTOTICGCCG GAATIGGTICGG GCCCTAGAAG AGCTICATICAG AGAACTICACC

CCATAGITISC AGCAGCAGCT GAGATCCTGC AGTGCTTTGG CTATGTCGAC

GATCAGCTAC COTOGTACCA TCATGGACAC CACAGTGGAT

TICCESCISC ISCAGISSET SCECEGGAAG SCCIACCISC SGAAISCIST SEISGIGAIC TICATCACCI CCACAGCCAT CCIGCCCCIG CIGITCGGCT GCCIGGGCGT CITCGGCCIC ITGATIGACT ATGGICICIC CGGCTACCAG GAAGAGICIG CCGAAGIGAA GGCCAIGGAC

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CCTCAGGGCT GGGCAAAGAA TGTGCAAAAG TCTTCTATGC TGCGGGTGCT

8 AGICATICIT

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GCTTGACCTC CAAGTAGAGC TGATACAGAG ATCTGTGAAT ATTGTGATAG

AMATICTITG TCCCAAGCCT

TOOCTITIGIT CAATGCTTCC ACTCTAGGGC AGGCAGAGCA GTCTATACTC

GTATTCATAC ATTTCAGCTG CAAGTCAGCA ATTTCCCCAGG TACCATGTAA GCTATAAAAC AAAGACAGAG GATAGCTGTG ACTICATOGGA TICATGAGGTIC

TOCAGGITCC CTITITICCTT CCTCAGGITT IGICICITEC IGIGITATEC CCAGCAAGGG CATGGCTGGT

AGAGACTOTO GOGTOGATIG GGAGAACAGA TTAGGAGTAT AGCAAATGAA CCCAGAATGG

> 1380 1320 1260 1200 1140 0801 1020

AACAGTGGGG GIGAATOCTG TATTOGCACA GGGANTAANT ATCCTGGCGT CTGGAGCCTT CACCTCTCCG AGCTRACTOT GRATGAGGAG AGTACCTGCT GCAGGACCTG GAGGTCAGGT

> 1500 1440

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GGGCATTGTC TGGGCTGCAG GGCTGCCAGG TTCTGTACTT GTGTCCAGCT CCTGTGATAC TGCCATGGCA CAGGATCTGA GTTGCAGCTC TGCACCCTAA

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GOCCIOTOTA GAGOCCCCIC CIGIOCCCIC AGIGGCIGIC GITTGITAAC AICAICAGGA GIGGECETGG ATOCTGGAGE TGGAGGGTTT TETGTGETEA GACTGTAGEE TGTAGETETT AGATGOGAAA GOTCAGOCAG AATTTTTCTG CCCTACAAAG GOTGGAAGAG AAAGGACACA 1680 1620

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CCTTTTGTCT CCTAAAGATA GGGATCTACT TTTGAAGGGA ATTGTTCCTC CCAAATAAAT GTCCCAAAGA AGAGCATGGG TGGCAGATGG TAGGGAATTG AACTGGCCTG TGCAATGGGC TITACCCICT AGCIGITITIA CITAGAATGI AACATATGCI GCCTACCCAC CICAAAATGI TCACATTIOG CCAAACCIGA CACIGICITG CIGCATICIC CITIGGCAAA CAICAGGGIC TIGOTITIACO TIGGIOCOTIT CITTIGIGCO AGIATICAAG IGGIATAGOT CIGAGOAGG GCAGAATGGG GTTGGGCCTG TGGCCCCCAA ACTAGGGGGT GTGGGTTCAT CACAGTGTTG TCACCTITAT TOTTGAAACT GAGGTTTACC TGGATCTGGC TACTGAGGCT AGAGCCCACA AGAACATOOT CTTCTTGCCA CGGGGTGTTG TTGTCTCTGG TGGTGCTGCA TGTCTGTGGC ATOGRACIACA AGOGOTICACA OCATOCCTICC TOCCTTACCO TOGCAGTACO GAGACAGTICC CTGTACTGCA AGAGGGCCCT GGGCCTCTGC TTTCCATATT CACGTTTGGC CAGAGTTGTA AGAATTCAGG ATAGCCCTTC CTAGGGCACT GGACTTTCTG GCATGGGGGC TGTGTTTGCA 60 540 480 360 300 240 180 720 660 420 120

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369		370
AATOTCATCT GCTCAAAGTT GAGTGGTTTA TTCACAATAA ACTOTAAGTT TCTGAITATA	1920	(B) TYPE: nucleic acid (C) STRANDENESS: double
Алааалаала аалалаала лаас	1944	(D) TOPOLOGY: linear
	,	5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 146:
		GGCACGAGGA GGGCCACGGC AGCCATCGCG CTTTGCAGTT CGGTCTCCTG GTGTACGGCC
(2) INFORMATION FOR SEQ ID NO: 145:		AACGCCAAGT AGGGGATTGC GTTCCCTCCA GTCGCAGACC CTATCAGATT TGGATATGTC
(i) SEQUENCE CHARACTERISTICS: (A) LENOTH: 1021 base pairs	01	CITCATAIT GAITGGAITT ACAGTGGTIT CAGCAGTGG CTACAGTTIT TAGGAITATA
(B) TYPR: nucleic acid (C) STRANDEDNESS: double		THAGHAACT GOTAAACTGG TATTTCTTGG ATTGGATAAT GCAGGAAAA CAACATTGCT
(D) TOPOLOGY: Linear	15	ACACATGCTA AAAGATGACA GACTTGGACA ACATGTCCCA ACATTACATC CCACTTCCGA
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 145:		AGAACTGACC ATTGCTGGCA TGACGTTTAC AACTTTTGAT CTGGGTGGAC ATGTTCAAGC
TICARCICACG COTICCGGGGT GOGGNACGGG GAOTTICCGGC TIGARACCCG TGCTCTGGGC	09	_
COGCOCCTIC ACCATGOCCT COGCAGAGCT GGACTACACC ATCGAGATCC COGATCAGCC	120 20	
CTGCTGGAGC CAGAAGAACA GCCCCAGCCC AGGTGGGAAG GAGGCAGAAA CTCGGCAGCC	180	AACCAITGCT AATGTGCCTA TACTGAITCT TGGGAATAAG ATGGACAGAC CTGAAGCCAI
TOTGGTGATT CTYTTGGGCT GGGGTGGCTG CAAGGACAAG AACCTTGCCA AGTACAGTGC	240 25	
CATCTACCAC AAAAGGGGCT GCATCGTAAT CCGATACACA GCCCCGTGGC ACATGGTCTT	300	
CITCICCGAG TCACTGGGIA TCCCTTCACT TCGTGTTTTG GCCCAGAAGC TGCTGGAGCT	360	_
ACTUTITIBAT TATICAGATITO AGAAGGAGCC CCTGCTTCTTC CATGICTTCA GCAACGGTGG	420 30	
COTCATOCTG TACCGCTACG TGCTGGAGCT CCTGCAGACC CGTGGCTTCT GCCGCCTGCG	480	TAACTIGAAT TCAATAGACT TITOCTOSTT ATAAAACAGA TGTTTTTTAG ATTATTAATA
TOTOGRAGIC ACCATUTITO ACACCICUC TIGATGACIAC AACCTIGITAG GGCCTUTACO	540 35	_
авсествося вселество лесоссавое соселеств состатис тествопос	009	AIGITAGCIT ICTAATICCA TAAAAGTACT TGGTTITTIAC AGTTIATAAT CIGACAICAC
CTTIGGCCTG GTGGTGGTCC TGTTCCAGGT CCTGCTTGCT CCCATCACAG CCATCTTCCA	099	CCCAGCGCA TITIGIAAAGA GCAACITICC AGCAGIACAT TIGAAGCACT TITIAACAAC
CACCCACTTC TATGACAGGC TACAGGACGC GGGCTCTCGC TGGCCCGAGC TCTACCTCTA	720	) ATGADACTAT ADACCATATT TADADGCTCA TCATOTTADA TTTTTTATOT ACTTTTCTGG
CTCCHGGGCT GACGAAGTAG TCCTGGCCAG AGACATAGAA CGCATGGTGG AGGCAGGCCT	780	AACTAGTTTT TAAATTTTAG AITHANGIC CACCTAICKT AAGIGTACAG TTAATAATTA
GOCAGGCCGG GTCCTGGGGC GTTCTGTGGA TTTCGTGTCA TCTGCACAGG TCAGCCACCT	840 45	GCTTATTCAA TGATTGCATG ATGCCTTACA GTTTTCAATA ACTTTTTTTC TTATGCAAAC
COSTGACTAC CCTACTTACT ACACAAGCCT CTGTGTGGAC TTCATGGGCA ACTGGGTCGG	006	GTCHTGCAAT AAAACAAACT CTAATGTTTTG GCAAAAAAA AAAAAAAAA NTCCAGGGGGG
CHGCTGAGGC CATTGCTCCA TCTCACCTCT GCTCCAGAAA TAAATGCCTG ACACCTCCCC	096	GENTISTATE CARTETION TABAG
ACHADADADA ADADADADA ACTOGROGOG GOCCOGOTA CCCADITICOC CCTNIDADGO	1020 50	
	1021	
	\$\$	(2) INFORMATION FOR SEQ ID NO: 147:
	1	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1346 hass nairs
(2) INFORMATION FOR SEQ ID NO: 146:		(B) TYPE: nucleic acid
(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1285 base pairs	09	(C) STRANDENKESS: double . (D) TOPOLOGY: linear
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0 € € TYPE: nucleic acid STRANDEDNESS: double

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(2) INFORMATION FOR SEQ ID NO: 148:

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Ξ SEQUENCE CHARACTERISTICS: LENGTH: 2098 base pairs

2 8 AAAAAA GACCTOTTOG CCTOGOGOCA CATGOGATOC TTCTAGOAAC ACAGTITGAG AACCACCAAA ATTTTRAAA TGAAAAAAA GCTGCTCTGA TTCAGGGGAT GTGGGTCGGG GTAGAACCTG 1380 1320 1386

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CICITICCAT OGIOGIOCCI OCICAICITO CIGAIGCAAA CIAGGAAGII AGOCIGCAIC

1560 1500

1680 1620

ICOGAJOTOGO TITICOCTOGA GAGOTOCTITI OCTOTOTOTO AGACTOAGIC ACTOTOTICO

GECTOACCTC COTTTCTGGG GAATGTGGCT GAGCTGTGGT AACCAGCTAC ACCCCAGGTG

CICCATCIGG TCACIGCAGG TGCCAACCCT TCATCCCCCA TGTTTTCCTG GGCCATGGAG

45

TRACTOTION CETANAACET AGETGGGCTG GTCTTGCTCC CAGCTTGCTT CCCCCCCCC

GUAROTOCCT TIGUAGUCC CIGIOGITICC TOCGUCACCA GIGIOCOTIGU CIGUCAIGUC

AAGCTCATCA GOOGCTTOTA CCCTGOTCAC CAAGCATGOT AGCAGCTGCC TGCAFTGTAT

1380 1320 1260 1200 1140

1440

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CCTGGCTGAG ACTGTCAGCC TCCTGGAGGA GTGGGGTCCA CCTTCTTCTT GCCCTATGCA

ACCITITATICE ATCHARGETE ACCATAGECE COTTECTION GACCIGGACE CICCATIGNA

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960 900 840 780 720

GIGCAAGCTT CACTICICAC CCAGCAAGGT IGACICATCT GCCICCATGT CICIGGGGCT

6 TOOTGGTGAA COTTGGACCA CAGCATGTCA GTGCTCTAGG GATTGTCTAC TOCAGGGATT ITICITICAAAA TITITTAAACA TOGGAAGITIC AAACAAATAT AATGTOTGAA ACAGATCAAA 1260 1200

AGCACCTICA TCACCAGAGG CITGAAGGAA CCCCGCCATG TGGCAGGGCA CAGGCACTGI GANGCCAGGG TOGCCAGCAG TOGCTOGGTC CTTCCGCCCC CCTCCGTCCT CCTTTCCCTG 1080

TOCHGOCCOT GOTGAGGAGG GOCGCCTCCC CAGAGTCTGC TCCCCCGTGA ACACAGAGCA 1020

TOTOCCAGGO GGTCCAGCTG GAGATGCAAG GCCCCTGAAG GCGCAGGCTN CCAGNCCGCC 960

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ACATOGTECT GGAGAAGGAC GAGACTOTOG ATGTOGTCAA CGGGCTACTG CAGCACATCC

AMATTGGCTT CCTGAATGAC AAGGTGGAGG AGCGGCGGGA NCGCTACATG GACTCCTATG

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840 780 720

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GTCAATCCAG AACTGCCTCT GAGCTCCAGG CTGACCACAG ATCAGCCACA GCCTGATGCC CACTECCTOG GGACTATIGGG ATCACTGTCC CCCCACCTGT GTGGCCACAC CATGTGTCCT AGCATGAGCC ATGIGCTICT TIGCCCTICT CIGICCIGIT CCAATCTICT GCCTCCCAGT CTTCTTCCCA CCTCAAGATG AGGTCCTCGC CCCCTTGTCT TGGCATAAAA ACACCTTTAA GOCCTICCCT GICCACIGGG CIGGATICAT GITCANACCA CIGGACIGGC AGGGCAACGA TOTOCCATICO TOCCTTOTICO CCAGCTICTO CAGGATGCCC TCCCTACCCA MCTYTYCCTG GCTGGTAACT TIGGCGCCIC CGCCAAGCCC TGCCAGACTC CCCTGGCTGT GATGGCATTC CTICTOTOGG TCTACACAAG AAGGTTCCTG TGAGGGCTAT CAGTTGTTGC CTTCTAGCTT

TOCHOCICA CITTOCICAC CCITCCCCIC CCCTCCICCI TCCTTCCACA CAGCAAGCCT

GAGACTETAT COGGGACETE ACCATOGOCG ATGGGGTTCC TGGTGTGCAG AACATTETCA TOTOTOHGAA CISTGOTIAC TICCAGCAAC TIGAGGGCAA AACCAATOTC ATCCIGCIGG

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CCAGGGATTT AAGGGCCAGC TGATACACAC ATACAACAAG AACAGCTCTG

9 60 540

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660 9 540 480 420 360 300 240 180 120

AGATGAAAGT GTTCCACCCC AACATCCACA TOSTSTOTAA CTACATOGAT TTTAATGAAG

ACATTECCCT TITCATCTIT TCTGCGGGCA TTGGTGATAT CCTGGAAGAA ATTATCCGAC

AGTCCANTGC ANTGCTCAGG GAGGGATATA AGACCTTCTT CAACACACTC TACCATAACA

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COCACAATOT CCTATOTOAG CAGAAGATTO AGAAGTTTCA GATAGCOCAG GTOGTTAGAG 480 420

TOTAL COGGACCOTO ANGUNGANGO TACCTONTAT GOTOGRATIGG TOTALCHANG

360

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CTTGGCCTAG CCACTGGGCT GGGATCTTCT GGGTCATGTG ACTGTGTATC CAGGAGCAGA

CCTTCRATGA AGTGAGACCC TTGGGGAGAA CGGGCTGTGG ATGAAGGAGT GGACTGCAGC CTGGCCATGG TTGCTGAGCA TGGGCAGACC AGTGGAGGCC ACCCTACTOT GTTATCTGCG

GACCTAGAAA TGGGCCTGTC TGGCATTTCA GAGTCAGGCA AAGCAGGCAG GGCCAGGGAG

AACTIGIATI CICAGGATIC AGGAICIACC CAGCACCAAA GAIGIATITI CAGGAGAACA

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AGENCETICITE COCCOCCOTTG GGACTICTGAC ATCTTAAGGC TGCACGGTCG TGTCCTTGTC

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 148:

(D) TOPOLOGY: linear

TOGGTGAGGE CATOTETOTO ATCCAAGGTT CETGGAACTG ACACAGGAAG GGGCTGTGAA

CCCTAAGTGG GTGTMATCTC CTCCRACCGA GGCTTCTMAC CCTGGAGATG GCAGTTACTC

15

TTOCATATAA TOGAAAGCGA TGCCCTTCTT CTTACAATAT TCTGGATAAT AGCAAGATCA TCAGTOAGGA GTGTCGGAAA GAGCTCACAG CGCTCCTTCA CCACTATTAC CCAATTGAGA 300

5

GCAAGGGCGS COGAGACCGG TTACAGGTGA TTTCTGATTT TRACATGACC TTGAGCAGGT 240 180

ATGAAGGCCA CGGTCCTGAT GCGGCACCTG GGCGGGTGCA GGAGATCGTG GGCGCCCTCC 120 GOCACGAGOT GOCOCAGGO TCAGTOOTTC TCTCGGOTCT CGOGACAGGT GAGCACCCTG (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 147:

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**GCGTCAGAGC** 

CTCCCCTCC CATTICAAGT 440 200 260

CACCACCTAC

620 680 740 8

GATGACCOTT CATTAATAAA TITIOCATCTC ATGCACACCA GITACTTCCT CITIGICATG TITIGECCEA GAAGIEGET TEGETEAAG CETAATTECA CAGETECTIG GCCTGTAACT GATTTGTACA TATTTATAAA AATCTATTCA GAAATTGGTC CAATAATGCA CTGATGGGAC GITTCCACGT GTCTCTAGAG AAGGATTCCT GGATCTAGCT GGTCACGACG FITTGETTTG TITTGETTT TITGETTTT GITTTCCTTT ATGAGAAA AATAAATAG ATCITITICAC CAAGOTCACA GGAGCATTGC GTCGCTGATG GGGTTGAAGT TTCGTTTGGT TCTTGTTTCA GCCCAATATG TAGAGAACAT TTGAAACAGT CTGCACCTTT GATACGGTAT TOCATTICCA AAGCCACCAA TOCATTITIOT GGATTITIATG TOTOTIGIGG TTAATAATCA TAGTAACAAC AATAATACCT TTTTCTCCAT TTTGCTTGCA GGNAACATAC CTTAAGTTTT GACGCTGACG AGAGAAGGCC TCTTCCTTGA GGGTTGGTGC TGTGTTGCAG TGACCGTGGC CAACGCTOTT CAGCAGGAAG TAAAATCTCT TTGTCGCTTG GAAGCCTCTC AGGTTCCTGC AGAGGAAGCT CTTTCTGGGG CTGGTGAGCC CTGTGACATC ATGGACAGCA GTGATGAGAT GGATGCCCAG GAGGAAAGCA TCCATGAGAG AACTGTCTCC AGAAAAAAGA AAAGCAAGAG ACACAAAGAA GAACTGGACG GGGCTGGAGG AGAAGAGTAT CCCATGGATA TTTGGCTATT GCTGGCCTCC TATATCCGTC CTGAGGACAT TGTGAATTTT TCCCTGATTT GTAAGAATGC CIGGACTISTC ACTIGCACTS CIGCCTTTTIS GACCAGGITIS TACCGAAGCA CTACACGCTG GATECTICCC TECCTITISCS TCTECEACCA GAGTCAATGS AGAAGCTGCS CTGTCTCCGS GRATTACGCC AACTCGGATC CGGCGGTCOT GAGGTCTGGA CGAGTCAAGA AAGCCGTAGG TOTTITIOCIA TOCTOTIATO AAGOGCAGAC CTAGCAAATT ITITITICAGA GAGACTICAGA GAACCATAAT CCTTGCCTGC TGAACCCAGC ATCCTCTGTG AATACATTAT CITGCGATGT TGGGTTATTC CAGCCAAAGA COTOCITITOC CCTOGGIACA GCCAGAGCCC TICAACCCCA CCITGGACIT TCACATTITIA ATACTACCAA AAAATGGACA AAAAAAGTCG AGGGGG (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 150: 374 (A) LENGTH: 1569 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 150: GTCATAACAA AATCCTGAAT WO 98/39448 S 9 15 2 22 ಜ 33 6 45 S 55 8 8 960 1020 1740 0081 1860 1920 1980 2040 2098 S 130 180 240 8 360 420 480 240 909 99 720 780 840 ACTOCCAGAG ATGCCAACAC AAAATACGCC CAAGATTACA ATCCTTTCTG GTGTTATAAG OSSCCATTG GAAAGTCTA TCATGCTTTA AATCCCAAGC TTACAGTGAT TGTTCCAGAT OCTOGRAPTIC CITITORGOT TOCTCTTCTG CITCTCACTC TIGIATIANG ANGGATTCCA INCACCACG COTECGAACT GAGGGGGGG CGGGAGCCGG TTGGKGTCTG GTCTTCGCGT COCITICAATO ATTITOGACTT GIOTOTATCA GAGAATGAAA CCCTCAAGCA TCTCACAAAC CCCCAGGCCC TOGAGGACTC GGGCCCGGTG AATATCTCAG TCTCAATCAC CCTAACCCTG GACCCACTGA AACCCTTCGG AGGGTATTCC CGCAACGTCA CCCATCTGTA CTCAACCATC ACCCIGCCIA CAGCGIGGAG CICAGAIGAC IGCGCCCICC ACGGICACIG IGAGCAGGIG GIATTICACAG CCTGCATGAC CCTCACGGCC AGCCCTGGGG TGTTCCCCGT CACTGTACAG CCACCOCACT GTGTTCCTGA CACGTACAGC AACGCCACGC TCTGGTACAA GATCTTCACA GOSTIACCIA CICTOCCAAG TGAGGACAAA CTGCTAGGCT GTATCCCATA ATTICAGGAT GAGAAACATT AACAATAAAA ATTTGTAGTA AACATAACCT CATGANGACT AAAAAAAAA CASCECECOSOS GACCAGACISE TISCECECOSOS GOGGGAGAA GATGOTIGECER AGCOGCETICO GOCCCOGOTO GOCCCGGGGC GCGCANTGCC CGCCGGGGCG GGGTGGAGCT GATCAGAATA ATGITICAGCA TCAACCCCCT GGAGAACCTG AAGGTGTACA TCAGCAGTCG GCCTCCCCTG GIGGICTICA TGAICAGCGI AANGCCCAIG GCCAIAGCIT ICCICACCCI GGGCIACTIC TICAAAATCA AGGAGATTAA ATCCCCAGAA ATGGCAGAGG ATTGGAATAC TTTTCTGCTA GACACCACAA CTCCCGGAAAG TACAATGACC AGCGGGCAGG CCCGAGCTTC CACCCAGTCC THAGGGCATC AGATTGGACT TTCAGGCAGG GAAGCCCACG AGGAGATAAA CATCACCTTC CAAAGGGAGA GTGGCATCCC TGCTGCTGCT GTGCCAGACC AGAGTTTCCT GAGGGGCCCT GACCCTAACC CTCCAGCTCA GCCCTGTACA CCTGACCCTG TAAATGAGTG GGGTTTGCTG STCACAGCCA GOCCOSCOAC GCGCCGCCAC GAGTGAGCCC AGCGCGACCG CCGGCGTCCG CCGAGCAGCT AAAAACTYOG GGGGGGCCC GTAACCCATT GGGCCCTTNO GGGGGWGTT TTAAAATT ACTIGIDATICS CITCACCAG TAAAACCAAA AGGACTICTIG GGGGTICAGT SEQUENCE DESCRIPTION: SEQ ID NO: 149: (A) LENGTH: 1847 base pairs (B) TYPE: nucleic acid (C) STRANDEDMESS: double (D) TOPOLOGY: linear 373 SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 149: (X 3 6 20 25 8 <del></del> 45 S 9 15 2 25 3 35

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PCT/US98/04493

375

CCCACGOGTC COGRAGGATT GACCAGTTAA CCAACATCTT AGCCCCCATO GCTOTTOGCC CTGTGAAAGC TGGTCTTAAA GAAGAGGAAA CTGAATTGAA ACAGCTGAAT TTACACAAAG TATICIATOTO COTODAGTAC GICCIGCICT GGAAGGITTA CCAGAAAAACC CCAGCICTAG AGATTATGAC ATTTOOCTCC CCAGICATCO OCTOTOGCTT TATTTCGOGA TOGAACTTOG SEQUENCE DESCRIPTION: SEQ ID NO: 151: TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 151:

35

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ATCAGAGGAA CCTTAGAGGC CTGAAATTOT TGCTTCCAGT TTAGCTOCCC CTCAAATTCA GTAGGGAAAA ATTAAACTCT TIGAATCTCC AAACAAGGAA GITTCAGCAT TCCCTTATGG TITICGIGCTA TIGAGITIGT TCTTTATICT TITATCCCAG TGAAAATIGT TGATCTIGCT ATTITAATOT AAAAATTIGC ATTIAAAAGG AGTGGCCCTG TITTICTGTGT TAAAAACCCCA OTTAGOGGAG GGAGAAATOT TGAATCAAGA GOGAAAACAA CTACTATGAT TTATAAACAT

AGTGAATATT TICCCTICIC CCTITACCCT ICICCAGAAA TAAAGCAGGI GACAGGGITT

1560

1500 1440

1569

23

1320 1260 1200 1140 1080

20

GOSTGOCTICG ATTICTATIATIC TOSTTAGTAA TOTACATICCT CTTCAGGTTC TAGGGCTCCT CCCATCCCTT GGGGGCAGCC TCGAGTGTAG TCCATTAGTA ATCAGATTCC AGTTTGGACA 15

GOTCOGARAC TOCOCAGTOR ACAGOGTOTO CAROTCATCC TOGACCCAGT OCACAGOGTT

1020 960 900 840

AGCACGGACA TGCGGCATCA TCGAGTGAGA CTGGTGTTCC AAGATTCCCC TGTCCATGGT AATTTICATICT TTATTICCGAT TOTICATOGGA ATGATATITA CTCTGTTTAC TATICAATGTG TACGAAGATG TICATACCAA ICCAGACCAG GACIGCIGCC TACTGCAGGT CACCACCCTC

COGCTCTTTG ACTOGTOGCA TCCTCAGTAC CCATTCTCCC TGAGAGCGTA GTTACTGCTT

0

S

TGGTGCAGAA AGATTOTTGG GAACAGACAG GAACCAATGT GGCAATTCAA CTTCAAGTTC

720 660

CTCAGGGACT GAGTGGGTTC CATCCTCAGT ATTTTGATGG GAGCATCAGC TATAACTGGA TIGCTTICCT TIATAIGACT GICCIGGGCT TIGACIGCAT CACCACAGG TACGCCTACA CCTTCCGAGA TGGATGGGTC TCCTACTACA ACCAGCCTGT GTTTCTGGCT GGCATGGGTC

ATAATOOGAA CTGTAGCTTT TACTTGGCTA CGTCGAAAAT GTGGTTTTGGT TCGGCAGGTC

600

480 420

600

780

AATOCAGOCA TIOCAGAAAG CACOOCAGO ACATIAAAGA AITOCAAAIG CITACITITO GCTTGTGTGA TOCGATCTCT GTACCATATG TATGAGCCAT TTGCTGCTCG AATCTCCAAG

AAAAAACAGT CCCCTAGGTT AAAGAGCAAG TOTACAGGAG GATTGCAGCC TCCCGTTCAG

8 E SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1540 base pairs

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ATGAGCTTGA ACATGAGCAA GAGCCTACTT GTGCCTCCCA GATGGCTGAG CCCTTCCGTA ATACTGAGCC AAAACCCCTG GAGGGAACTC ATCTAATGGG TGTGAAAGAC TCTAACATCC 240 .00

> ઝ 30 23 20 2 5 GTGTACAGAA CICCATGAAC TATCTICTIG AICTICIGCA TITICATCATG GICATCCIGG TRACTOTICAC ACACTTOCTO CAAGAAAATO TRATTIGAATO TGAAAGAGGO ATTATAAATG AUGGICIAA TICIGCIAAT ATIGICCCGG AAGGAGAGTC AATTACACCT ACCAAGATAC CTGAAATTAC AACTGAAATA TACATGTCTA CTGGAAGCCC CCTGGACTTG TCCGTTTCTC CTTTTGAAGA TATCCGATCA TGATETICAGO ATTROCAGAS CTITICETGTT TGATETTGTG TGTGATETET GTATTCATGE OTTTAACTGT OTCCTGATGC AAAAGAAGTT AGGAAGGAAA ATCAAGCAAA TACATCTGTT GTTTGAGACA GCCACATTAT GTATITICCGA TITIGCCCAAA ATACTCTGGG AAACAAGCTC TITIGCTTGCG CTCCAANTCC TGAAGCTTTT GGCTTGCTCG TATTGATTTC AGTCTCCTTT GTGGCAATGG CIGICAGICI CAGAATTOCA ATAAATGGCT GGGTGTTTTG CTCTGTTTTT ACCACAGCTG TGCCTTGAGA ATAATOGATA ACACTGRATT CCCCTATTTC TCATGAGTAG ATACAATCTT ACGTAAAAGA TTGAGGGCCA TOGRAAAAA ATTGGGAAAA GGAAAAACTC AGTTTTAAAT ACGGGAGACT GCTGTTTGCA GITTAGGAAA CCTAAGTCAG CAGAAATTAA CTGGATTAAT TTCCCTTATG TOCTATION THACTAGATT ATATAGAGCA CATGROCTEA TITTIGHACTG GOCGICATIG CIGCTAGAAT COGICTITIGG TCCTTTGATT AGACAAGTCC TGAATCTGTG CCCATAATCT 1200 1140 1500 1440 1380 1320 1260 1080 1020 960 900 840 780 720 660

25 (2) INFORMATION FOR SEQ ID NO: 152:

8

GIGGITAGIC ACGIGAATIC AGITATCATT IGACAGATIC

1540

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1719 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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ž SEQUENCE DESCRIPTION: SEQ ID NO: 152:

SS

TACTTATGAG GTCAATTGGA AATAAGAACA CCATTTTACT GGGTCTAGGA TTTCAAATAT TAGCAGECAT GICTAGEATE ACCITICETG CIGICAGIGE ACTIGITICA CGAACIGETG TACAGINGGC ANGGIANGGC TINGGINCAG AACCINGGAT GANGNGGGCT GCNGGGGCAG 180 120 8

8

376

PCT/US98/04493	
WO 98/39448	

(2) INFORMATION FOR SEQ ID NO: 153:

(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 861 base pairs
(B) TYPE: nucleic acid

WO 98/39448

PCT/US98/04493

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 153:  GCCACCAGGG AAGCCGGAAC ACTCTCATT CAAGGAACTA AGAACCTCT CCCTCAGGG TAGGGAGACA GCCAGGAACG AGTTCCTATT CAAGGAACT AGAACCTCT CCCTCAGGG TAGGGAGACA GCCAGGAGG GTTTTCTGGA AACTGTGGA TGTGCCTTT GGGCCCGG AAAACAGAAG GAAGATGCT CAGACAGTA ACTACACCT GGTGCTCTCT CTGCAGTTCC TCCTCCTCTCT CATCACCTCT TTGTCAATT CCTTCTCAGA ACTGCTCTCT CTGCAGTTCC TCATCCTCAT TGTGCACT TATCACACT ACTACACACA ACTGCTCAA AAGATGGAC TTTTCCTCAT TGTGCACTCT ATCATCCAAA ACTCCAACA ACTGCTCAAC ATCACCCTC ATAAGTTCAA AGGGACCATC ATCATCAACA ACTCCAACA CCTTCATATTGG ACAGATGGAC TTTCAAATGCT CATTCTAATTC CAGAGACTAG CTGCATGTT GTACTGCTAC TTCTTATAAAC GGACAGCCGT AAGACTAGGC GATCCTCACT TCTACCCAGA CTGCTTTGTGG ACAGATGGAC TTCAAATGCT CTTTCTAATTC CAGAGACTAG CACACTGAT GAATTGTGG ACGACAGAG TTCAAATGCT CTTTCTAATTC CAGAGACTAG CACACTGAT GAATTGTGG CTGCCTCAGA TAGAGAGGAC CATTTCTAACG GATCCTCTTT TCTACACGAA CATTTCCAAGG AAGTTCAACAC AGTTCCAAAG GGATCTAGA TAAGGCATGT TGTTTCAGCC ATTTCCAAGG AAGTTCAACAC AGTTCCATAGA TTAGGCATTT TGTTTCAGCC ATTTCCAAGG AAGGGGAATTT TCCAAGGGAA GGATTCAGAA TTAGGCATTT TGTTTCAGCC ATTTCCAAGG AAGGGGAATTT TCCAAGGGAA GGATTCCATCA TTAGGCATTT TGTTTCAGCC ATTTCCAAGG AAGGGGAATTT TCCAAGGGAA GGATTCCCTTC CTT (1) SEQUENCE CHARACTERISTICS: (1) LENGTH: 1101 base palis (2) TTTPS: nucleic acid (2) TTPS: nucleic acid (3) TTPS: nucleic acid (4) TTPS: nucleic acid (7) TOPOLOGY: 11neax (A1) SEQUENCE DESCRIPTION: SEQ ID NO: 154: (A1) SEQUENCE DESCRIPTION: SEQ ID NO: 154:

TIGITACTAG GIGCIGGAGG AACATCCCAG TICACAAAGC CCCCATCTCT TCCTCTGGAG AGGICACATG ACACAGGCAG AGCAGAGTGG CACCCACCAC AGAATACAGT GTGTCTTATT CCAGAGCCTG COGTOGAATC AAGTCCAACT GAAACATCAG AACAAATAAG AGAGAAATAA GAATAGAATG AATGACCCCA AAATARGOTT TTCTTGGGCG AGGATGTGCT GGATTAGGAA ACCIDENACE ACCIPITANG CETANGOTEE TICTACETAE CTOSTATIOS CATTICAGGT 

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380

25 6 35 30 25 20 8 SS 50 CACGGICCIC CIGGGIGCCI ACCAAGCIIG GITTIGIACAA AAGCAAGGIG GGAGICTATI CAATTAACCC GITTGAGGCC TAGGITGITT GGCAAGCCCC NGGCCTAAAG TITTAATTCG 2 CTCGTGCCGA ATTCCCTGCA G CITCATITAA ATACATCIGI GIGCATACAG ATACGCATAT AIGIGIGIGC GIAIGCATAT TAGCOTTTTT ATATATGACC TITGATTICT GITGITIGTA TITTAGCACA GIGTATGCAC GTTTACACTG ATGCCTTCCC TGCCCACCAC AAATTGTGTA CATAGTCTTC AGAATGATAC TICCTAGGGC TOGRAGGITT AGCAGCAGCC TOGTGCAGTG CCCTOTCATC AAGACAAACC GCAGAGCCAA GGGCCTGAAA GGAAGGGAAA GGGGAGGGTA GCGGGAAGGT AGCAGGTGAG TTCAGGTATA AAAACAAGAG ATTATAATAA AAAANTAAAA GAACCCTAAA AAAAAAAAAC ACTATOTOTO CTITITAGAA TAAAGAITAC ATATCATCAT TCCTITOOGG AAAATTOITA AGATGAGTAA TIGITATIGA AGATAGICAG IGATAACCAC IGACCAGAIG CTATCAATAC GACACCCIGC ATCCCIGCTA ATAGIGITITG CCACAAGIAT TAGIGAGICT TCCTTATIAA CACCCCTTTC CCCAGCTCCC AACCAAGAGC TOGTTCTAGG CCTGTGTTAT ATGTCATATT TITIGIACATG AGATACATCA CACTTACCTG TGGGCCAGTA TIGIGAAGTG AGTCTGAGTT ATCTCTCATC TGTAGTTTCC AAGAGTTCAG CTGAAGCAGA TGGAGTCCTG CAGCCCAGGI INFORMATION FOR SEQ ID NO: 155: Ε SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 2031 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 1101 1080 1020 960 660 540 480 420 360 300 240 180 120 8

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TGAAAACATG CCACCTCAAG GCTGGGCGCG GTGGCTCACA CCTGTTAATC CCAGCACTTT

GOGADOCCGA GOCGOGCOGA TCACCOGAGT COOGAGTTT GAGACCAGCC TOGACCAACA

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CAGGANAGGC TACCTARCTT CACATATCTG CAACCAGAGC AGCCACCAAG CATTACTTAG

GOOTAGAAAA TTCACAGTAG GAATGATTGT TAAGAGAGAG TGCTTGGAAC CATGGGTTAA

1320 1260 1200 1080 1020

900 840 780 720

CAGCAGGAAA ATGATTGTAT TIGAGTICCT GIGIGICCAA AACTGAGGCA CCAIGTICTI

CAGCCCAGGG CTTGTCTGGA TCAGCACCAA CGATTTTAAA GAAAAAAAGGA AGAGTTTCTT TGTATTTTGT TCTTGAGAGG GGTCAGTCTA GAAGCTAGAT CCTATCAGGA TGAGGAGCAG CAAGCCTGCT GAGTATTGCA GCTGCATTTG CCCAAAGGGA ATCCAGAACA AGTCCCTCCC ATCCAACTIT AATAGTATAC ATTIAAAAGA AAAAAAACAA AAGCCCTGGA AGNITGAGGC GTGATCCTCA GGTTGTGAGG TOTOTOCTTT CAAAAGGACT TITCTCTCCT AGCTGACTGA CTCCTTCCTT AGTTCAAGGA GAACTOCICAG CAAGCCTTGA COCCTTATGT ATGTAGCTGA GTCAGCAAGG TACATGATGC ACAGCTGAGA CAGACCTCTG CTGAGTAGCT CTGTGATGAC AAAGCCTTGG TTTAACTGAG COGAAACCCT CTACTOCCCC ATAAGCCAGG AAAAGTGAAA AGAGAACACA GTTCCTTTAA TITATTAGTC CCCAAGGCAA ACACAAATAT TAGATTAATA 900 840 780 660 600 540 720 480 420 23 20 2 5 S AMATCAMGGC ATCACAMTTT CTGTTCAGCG GGCAGGAMTA GGCTGTGAMT TGCTAGCACT GOCCCAGGOT GAGTGAGATG AGOTGCAGOT GOCTCATGGC CTTCTTAGAG CAGAGAGAGAG GICACICTIG CANGGCTICC ANGICIOGIT TOTGGCATTT GGGGATAAGT GCIGAACCAG GAGGAAGAAC TITIAGAAAC CAAAIGAICT TAATIGITAT IGCCCACCCC IGGCTITIICC TOCCAMATOT AGACTAGAAT CTOCAGGATO COACCOCACTO TATAOTTOTO CTTTCCCAGA CCAGAAACIG AGGCACAATA GOGTTATGAC TTGCTCAAGA ATATGTAGCT GCTAGGGGGT AGCATTIGCA GITTGITIGA GGCCICGITG CCAAIGATAG ATCACICCIG TIGACCIGGI AGIGAGGGIC TIOGGAIGAC TIOCIGIGIF CCICAAGCIG CACTITIGGG CCAICTCIGC TITITITIAA GCAATTACIT TIIGACITGI TOOTOIGAAA GTGCAAGAGG OGTACACCIT AGTATISTICAT TITTACTAAGT TECTAAACAA ACATTTATISC AGGCAACACT CETTIGCAGAT AUGICIOCIT OCTIOCIOCI TITICCITICCI TICICITIGGA AGAGGAAAGG ACTCIOGICA AGIATIAAGO COCCITITIG CITGGIGGIA CICIGICIGI GOCIGIGIGI GIGIGIGATA PATTITICATI TCAGAAGACT GAAGCAAAGC TGATAGTOTT TGCTGTTTCT TTGGCAGCTA

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2 INFORMATION FOR SEQ ID NO: 156:

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TATRATICTICA GCTACTTOGG AGGGYTGAGG CAGGRGAATT GCTTGAACCC RGGANGGCGG

TOGGRAAAAC CCCATCTCTA CCTAMAAATA CAAAATTAGC CGGGCGTGGT GGCATGCGCC

1800 1740 1680 1620 1560 1500 1440

1860

CAAAAAYICC GICTICAAMW MRIGCCGAAT ICGATATCAA GCTTATCGAT ACCGICGACC

1980 1920

2031

TIDAGITIDAD GAICOTOCCA TITOCACTICC GGGCCTTGGG GCAACAACAA

TOGRAGOGOGO GCCCGGTRCC CRATTCGCCC TRIRGNGRTC GTATTRCRAF C

Ξ i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1981 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

S

SEQUENCE DESCRIPTION: SEQ ID NO: 156:

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	WO 98/39448	PCT/US98/04493	. WO		_₽	PCT/US98/04493
	381			382		
	CCTGCACCCT GAGCCCTTCA CCCCTCCGAG TTCCCCCCAG GTTGGCTTCC TTCGATTCCT	09		ANTICTAITG TITTGAACTC TGAITTAAAA TIAAAITGCA GCTGGGGGTG GTGGCTCATG	1860	
	ITTICITIGGIA TCAACOTITIG ATTIGGAAGAA CAACCCCCTC TTTIGTCAACC TCAATAATGA	120		CTIGINATICS CARCATIAG GRACINAGGR GAATCACTIG ASCYCAGGAG TYCTAGACCA	1920	0
. 1	GCTCACTOTIG GAGGAGCACC TCGGGCACAG CTCACCGTYA TGGTCATTOT TACCCCCAA	180	'n	ATCTGGGCAA MAGAGAGC CCATCTCTT TAAATAAAA GTTAAATTGC TTAAAAAAA	1980	0
	GACCOCANAN ACTICTOTOTO GACACAGGAT GGACCOCTICAG CCCAGATICCT GCAGCAGCTT	240			1981	
2	GTGGTCCTGG CAGCTGAAGC CCTGCCCATG TTAGAGAAGC AGCTCATGGA TCCCCGGGGA	300	01			
3	CCTGGGGACA TCAGGACAGT GTTCCGGCCG CCCTTGGACA TTTAGGACGT GCTGATTCGC	360		(2) INFORMAȚION FOR SEQ ID NO: 157;		
	CTGTYTCCTC GCCATATCCC GCGGCACCGC AGGCTTGTGG ACTCGCCAGY TGCCTCCTTC	420	:	(i) SEQUENCE CHARACTERISTICS:		
15	TECCOGOGOC TECTCAGCCA GCCGGGGCCC TCATCCCTGA TGCCCGTGCT GGGTNATGAT	. 480	15			
	CCTNCTCAGC TCTATCTGAC GCAGCTCAGG GAGGCCTTTG GGGATCTGGC CCTTTTCTTC	540		(C) STRANDELNESS: double (D) TOPOLOGY: linear		
8	TATGACCAGC ATGGTGGAGA GGTGATTGGT GTCCTCTGGA AGCCCACCAG CTTCCAGCCG	009	20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 157:		
	CAGCOCTINCA AGGOCTICCAG CACAAAGGGG CGCATGGTGA TOTCTICGAGG TGGGGAGCTA	099	Ü	GAATTCGGCA CGAGCGCGC CATGGCGCTC CTGCTTTCGG TGCTGCGTGT ACTGCTGGCC	<u> </u>	
	GTAATGGTGC CCAATGTTGA AGCAATCCTG GAGGACTTTG CTGTGCTGGG TGAAGGCCTG	720	ž	GOCTITOTICS COCTOSTICS STITGCCCAAS CICTOGGAGS AGAICTOGGC TOCASTITICS	170	
25	STECAGACTO TGGAGGCCCG ANGTGAGAGG TGGACTGTGT GATCCCAGCT CTGGAGCAAG	780	_	GAGCOGATGA ATOCCCTOTT COTOCAGITT OCTORGOTOT TCCCOCTIGA GGTATITIGGC	18	
	CTGTAGACGO ACAGCAGGAC ATTGGACCTC TAGAGCAAGA TGTCAGTAGG ATGACCTCCA	840	-	TACCAGCCAG ATCCCCTGAA CTACCAAATA GCTGTGGGCT TTCTGGAACT GCTGGCTGGG	240	
30	COCTOCITIOS ACATGAATOC TOCATIGGÁG GOOTIGOTIGGO TGAACATGOT GAATCATOTO	006	30 1	TTOCTOCTOG TCATGGGCCC ACCOATOCTG CAAGAGATCA GTAACTTGTT CTTGATTCTG	  	
	CAACAAAACC CAGCCCCAAC TITICTICTIC ATGCTCCAGC ATTGGGGCAG GGGCATGGTG	096	U	CICATGATGG GGCTATCTT CACCTTGGCA GCTCTGAAAG AGTCACTAAG CACCTGTATC	e	
	GOCCATOTAG TCTCCTOGGC CTCACCATOC CAGAAGGGA GTGGGAGCCA GCTCAGAGAA	1020	۶۶ ر	CCAGCCATTG TCTGCCTGGG GTTCCTGCTG CTGCTGAATG TCGGCCAGCT CTTAGCCCAG	450	
35	GANCTEANC CCNGGNGNTC CNTCCACCTA TTAGCCCTOG GCCTGGACCT CCCTGGATT	1080		ACTANGANGG TGGTCAGACC CACTAGGAAG AAGACTCTAA GTACATTCAA GGAATCCTGG	480	
	TCCCACTCCT TTCTTAGTCT TCTTCCAGAA ACAGAGAAGG GGATGTGTGC CTGGGAGAGG	1140	∢	ANOTHGAGGA TCTCTGTCTC TTTARGCCAT GCAGCTGTCA CAGCAGGAAC ATGGTAGAAC	—.5. —.5.	
- 6	CICTISTICIC TICCIOCIOC CAGAACTIST OCCIAGACTT AGCATOCCCT TCACTGCAGT	1200	40 A	ACAGAGTETA TEXTETTETT ACCAGTATAA TATECAGGGT CAGCEAGTST TGAAAGAGAC	8_	
	STCAGGOCTT TAGATOGGAC CCAGGGAAAA TOTGGCCCTT CTGAGTCACA TCACCGACAC	1260	*	ATTITICICIA CCIGGCACIG CITICICITI TIAGCITIAC IACICITITIG IGAGGAGIAC	9 	٠
	TGAGCAOTIGG AAAGGGGCTA THIOTOTHIG AATHGACCAC ATTGAAGGAG CACAATGCCC	1320	¥ 4	atottatica tattaacatt cotcatotca tatgaaaata caaaataac agaaaagaaa	_2.	
45	TECTIFICITIG ATGCEACTTE CEAGGORGA GACAGTGGAA AAGAAGCGAG GACAGGAAAG	1380		TITIAATCAA CCAAAATTCI GAIGCCCCAA AIAACCACTT TIAATGCCTT GGIGTAAGIA	6-	
	GAITYGOTAG OTGAAGGGGT CAGSGGACTG GTAGTCACCC AATCTTGGAG AGGTGCAAAA	1440	Ę	TACCICIGAA CITITITICIG TGCCITIAAA CAGAIAIAIA ITITITITIWA ATGAAAATAA	8	
20	AGCACTOGGG GCTACCCGTT AGCTGCATCT GCCCTGGCTG TTTGCCCGTT CATGTCACAA	1500	50 A	AACCATATAT CCTATTTAT TICCTCCTTT TAAAACCTTA TAAACTATAA MAAAAAAAA	8.	
	ACTOCCACTA CTATGTACCT GCAGTGGGGT TGCAGAGATG GGGGAGACTC AAGTCTTACT	1560	æ	aaaaaaaaa ctcga	915	
	CCCCAGGAGC TCCCAGGGCC CAAGGAGGAG AATGCTGCCT CCTTTCAGTC TGGTCTACAC	1620	ť			
55	CCACTITICTS STAGCCICTC TECTICCIST ARTICISACT STITITICCAS ACTUASCICA	1680		٠		•
	AATAGTOCCC CTCCTTAAGC CCATCCCTCG CCCCCAGCCT GAGGTGATCT TTCCCTCCTC	1740	<b>.</b>	(2) INFORMATION FOR SEQ ID NO: 158:		
9	толаститта слеслетнас тотстеттся стесеттоо слесслела слетовсята	1800	09	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2117 base pairs		

WO 98/39448

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

Ĕ. SEQUENCE DESCRIPTION: SEQ ID NO:

2 5 S COTCAGCAAG ATCATCOCAG AGAATATTTA CGAGGGTGGT CTGAACAGTG ACTATGTCC TOGACAGCAT CTTGCAAAGA AAATCATCTT AAATGCCGTG TTTGGTTTCA TAAACAACCC actractractia acaccatica trastacaasa astraaasaac atchaccira aactraaccot COGGCAGAAG COGAGCCTTA GCCOGGAGGC ACTGCAGAAG GATCTOGACG ACAACCTCTT GOCCOOCOTO CTCACCOOCT ACATOTACOC OCOTOTOTAC TOCCTOTTOS CCGAGTOCTO NAROCCCARG ARACCTOTCA COCTOTOCOT GCACGGGTGG ACAGGCACCG GCARARATTT AGAGCGAAGC GAGGOTGGCG CGGOTCCGGG CATGAAGCTG GGCCGGGCCG TGCTGGGCCT 420 360 300 240 180 120

23 20 30 TGAAATGGAT AAGATGCATG CAGGCCTCAT AGATGCCATC AAGCCTTTCC TCGACTATTA CATCAAGCTC AAAGACATTG AACACGCGTT GTCTGTGTCG GTTTTCAATA ACAAGAACAG GITACAGITG TOGATICGAG GCAACGIGAG TOCCTOTOCG AGGICCATCT TCATATITGA CCTGTTTGTG GCCACATTGC ACTTTCCACA TGCTTCAAAC ATCACCTTGT ACAAGGATCA AGCAGAAAGG ATCACAGATG TGGCTTTTGGA TTTTCTGGAGG AGTGGAAAGC AGAGGGAAGJ TGACCTOGTG GATGGGGTCT CCTACCAGAA AGCCATGTTC ATATTTCTCA GCAATGCTGG 780 720 660 600 540 480

35 CCCCCTGGAA TACAAACACC TAAAAATGTG TATCCGAGTG GAAATGCAGT CCCGAGGCTI TOGOTTOTOG CACAGCAGOT TAATTGACOG GAACOTOATT GATTATTTTG TICCCTICCI 900 840

TGAAATTGAT GAAGACATTG TAAGCAGAGT GGCTGAGGAG ATGACATTTT TCCCCAAAGA

COATGATTGA CAGTCATGAT TOOCAGCCGG TICICAGATA AAGGCIGCAA AACGGIGIIC ACCAAGITAG AITATIACIA AGTCACTOCC TOGAGTTOGA ANAGANACAA 1020 960

6

CATGACIGGA CAGCAGCTIT TOTTICTIGA CCCTOTAATA TGACAGICTG CTAATATIGA

ATTAGCAGGA

CICITICITY CONTICITY CICCONTY CCCONCIGAC CCCAMACGIT ATTOTCCAMA CANAGECTIT ATTITICGEAG TIMAGECANA TETETITITEC AGNANGITAG TIMITITICIE TOGISTICAGE TOTOTITITAT TOCACACCTA AATCCTGATT ATAGGETITT CATTITCTCCG

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TOTTOCTOTT TICCACACIT CCACCCCCAG CICCITICCC TGGAAGAGGA ATCCACTGA 1140 1200 1080

GGGACCACOO GCGAGGACGT TGATOTGACA GGAATTCTCC GAAGCCTCCC GAACACGCAC AGAAGGAAGG CIGGCATIOT TICCACCCC TOGTGCCTOC 1260

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TOGAGCATGA GCCAGCTCTG TCCCAGGATG GTCCCAGCGG ATGCTGCCAG GGGCAKTGAA

1620 1560 1500 1440 1380 1320

8

CCTGTTTTCT AGGTGGAAAG AATCTGATGA GCCTTTTAGG CCTGTTCAGG CAAATTTGAG

900 840 780 720 93 600 540 480 420 360 300 240 180 120

CACACATOCA TOOCATTTOT TCCACAGGAG GGCATCCCTG GGGATGTGGC

TITMAGITICT CATCATTAIT ACATAGACIG TGATGTATCT

TITIDACATI AUGAGAGACT OCTCAGATIC TAAGITOTIG OCCITIGIGIG CIGITICAAG GAAGGATGAA TAAGITITIAT TGAAAATGIG GTAACITITAT CAGCCCACTC ATCGCAGGGC TCATGATTTT TTACAAATTA TGTTTTAATT

S

TGAGCCCAAG

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8 3 GAAGAAACAA GCTCTGAACA GACTACGTGC TCAGCTTAGA AAGAAAAAAAG AATCTCTAGC GGACAATTIC CAAAGGCTIT AAAAAAAKGA CITGITICIG TIGIACIIGA GICITAAGAA AAAGIGGCCC ATAGITIIAGI CAGAAGGIGC AGITTITIGGG TIATAGICGT GATTITICGCT AATCAATCAT AGTACCACCT GTATTTCAAA ATGGGGGACC CAAACTCCCG

TOACCAGITIT GACTICAAGA TOTATATIGC CITTOTATIC AAGGAGAAGA AGAAAAAAGTC TACTCAGGAG ATGGATTTCA TICTTIGGCC TCGGAATGAT ATTGAAAAAA TCGTCTGTCT GGATGTGGTA CAGCTCCATG CTCCTCGATA TCAGTCTATG AGAAGGGATG TAATTGGCTG CCTGANAGGT GTGCGAGATT CCAGCTATTC CTTGGAAAGT TCCCTAGAGC AGCACTITIT GAAGTGICIG AGGITATACC AGICATGACA AATAATTATG AAGAAAATAT TTTTACAGAA

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(B) TYPE: nucleic acid (A) LENGTH: 2395 base pairs

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SEQUENCE DESCRIPTION: SEQ ID NO: 159: (C) STRANDEDNESS: double (D) TOPOLOGY: linear

TOTTCCTTAA TCCCTTTTCT AAAAAGGGGG GAAAATCCCGG ATGGATTTTA GGGATTGGTC

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CACACIGICA ACATTIGITA GAACCAGICT TITGAAAGAA AAGTATTICC AACTIGICAC GTAACCAATT TGNCCCC ATAMAATTIT GCATGIGCAA AAMAMAAAAA ANAAMAAAA AAAATCCCGG GGGGGGCCG CONSCICTOR GOIGGATION AAATTANGIG AATTITIGCCA TATTAAANON ICCICATITIA 3 CTTCTGAGGA TOCTGAGAAC GGTGTCTTTC TTTATAAATG CAAATGGCTA CCGTTTTACA TACTATTATT TOTTACOTIC AATCAGAATC CCCGAAACCT CCTATAAAGC TTAGCTGCCC TIGCCAGICA CICCGITIIG CAMAGGIGG CCCTICACIG ICCATICCAA AIAGCCCACA AACAGOCTOT COAAGGGTTT TOACOTTAGO AACAATGGGA GOTGTGGGAG TGATTTTGGO INFORMATION FOR SEQ ID NO: 159: ε SEQUENCE CHARACTERISTICS: 2100 2040 2117 1980 1920 1860 1800 1740

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GTOTTTAGGT GAAGGACAAG TAGGTAAGAG GACGCCTTCA GGCACCACAG ATAAGCCTGA

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ITTCATCATG GIGACTATGA AAAACAGITTI CIGCATGTAC IGAGGGGCAA GGACAAGACT GGAATCGTTG TCAACAATCC TAACCAGTCA GTGTTTCTCT TCATTGACAG ACAGCACTTG CAGACTICCAA AAAACAAAGC TACAATCTTIC AAGTTATGCA GCATCTGCCT CTACCTGCCA CAGGAACAGC TCACCCACTG GGGCAGTTGG CACCATAGAG GRTCACCTCC GTCCTTATAT TITCHTGITT TCTTACCACT TTAITCTTTC AGAGTITAAA GAAAATGGAC TCATGCACAG AACACTATGC ATTITGAAAC ITGITCATCC IGGATTITIT TAAATCATTI ITAICTCAGA

GCTGCACTGC ATCAGTATCT TACTAAAAT GTGAAATGAA AGGACTATTG TACACTGAAA ACTTAAACAA AAATTAGATG TCGTGCACGG ACTGTGTGAA AGAAGATGCT TTGCATATTT

TECTTABATE TATCTEAAAG CACAAGGTGA TACTCATTTT TATGGTCTTC CCATTTGTGC IGGITITITIC CICTITICACA ICICICATCA GIATITIAGAG GGICAGAAGI GAAIGIAACA GGTATAAATA ACATTTTTAA AAACAATAAC TTTGCTATAA TCACAGTTGT TCCAGAGCAC

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CATGITHGIT ICCIGGIACT CAIGIAITIT ITITITICCAG AICTCITICC CCAAGIIGCT

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IGTCAGATAC ATTCTAATGA CCAGAACTÖG TTTAAAAAA GAAAATACAA CCATGGGAAA GAAATCTTAA ATGAAAAACG CATCTCATTG TAGGCATTTT TGCCTCATAT TTTACTGGGC

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CTGAAGTAGC TATAAAGCAG CTATAAAACA GAAATACATG CATAGCTGCA GAAACCATGA

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TAGGTAGAGG ACTITITCTIT 1GGITTIGIT ITGITTIGIT TIGITITIGIT TITGGITTITA

NITICTAAGAG TATTCTGCTG COTOTGGATG CAGTTATACA CATTAAAGCA GATCTGGAGT

GCCAGAGTAG AGTACTGACC AGCAAATGG AGAAGATCAG AGAATGCAGC AGCAGTTTTT

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 160:

2	CCCCGGATAC CGCCTGACGT AGTGCCAATC ACACCTCTCG CGTCTCGGCG CCTCGGAGGC	-
	TAATGAGGAC GCCTGGGGAA AGGCAGTAAC GGATTTCCGG GTGGACCTTC GCTTTACGGC	
2	TOSTERASTIC TICOSCICIAA CICAGAGGAA GIGGGAGAGC AGITTANGAC AGICCICGGTC	
2	GIGITTAGGG CGGCCCCCC TGCGCGCCCA TGTTTCCTCT TTTCCTGGTT TCTCAAGAGT	
	GCTGCTGCTA ACGCGGTCCC CGGCACGCAC CATCTGTTGC CATCCCGGCC GGCCGAGGCA	
15	TTSCAGATT TOGAGATGG CAAAGTTCAT GACACCOOTG ATCCAGGACA ACCCCTCAGG	
	CTGGGGTCCC TGTGCGGTTC CCGAGCAGTT TCGGGATATG CCCTACCAGC CGTTCAGCAA	
ç	AGGAGATCGG CTAGGAAAGG TTGCAGACTG GACAGGAGCC ACATACCAAG ATAAGAGGTA	
3	CACAMATRAG TACTCCTCTC AGTITIGGTIGG TGGANGTCAM TRIGCTTATT TCCATGAGGA	
	GGATGAAAGT AGCTTCCAGC TGGTGGATAC AGCGCGCACA CAGAAGAGGG CCTACCAGGG	
25	GAATCGAATG AGATTTGCCC AGAGGAACCT CCGCAGAGAC AAAGATCGTC GGAACATGTT	
	GCAGTTCAAC CTGCAGATCC TGCCTAAGAG TGCCAAACAG AAAGAGAGAG AAAGCATTCG	
ç	ACTGCAGAAA AACTTCCAGA AACAATTTGG GGTTAGGCAG AAATGGCATC AGAAATCACA	
2	GARACCCCCA GACTCTTCAG TTGARGTTCG TAGTCATTGG GAAGTGAAAG AGGAAATGGA	
	ITTICCICAG ITGATGAAGA TGCGCTACIT GGAAGTATCA GAGCCACAGG ACATTGAGTG	
35	ITICTOGGOCC CTAGAATACT ACGACAAAGC CTTTGACCOC ATCACCACGA GGAGTGAGAA	-
	GCCACTGCGG ASATINCAAGC GCATCTTCCA CACTGTCACC ACCACAGACG ACCCTGTCAT	
Ş	COSCINACTIG GCANANCTIC NOGGGANTICT GITTGCCHCT GNTGCCNTCC TGGCCHCGCT	-
}	GATGAGCTOT ACCCOCTCAG TOTATTCCTG GGATATTOTC GTCCAGAGAG TTGGGTCCAA	
	ACTETICITY GACAAGAGA ACAACTETGA CITTGACCTE CTGACAGTGA GTGAGACTGC	
45	CAATGAGCCC CCTCAAGATG AAGGTAATTC CTTCAATTCA CCCGGCAACC TGGCCATGGA	
	GOCARCCTAC ATCAACCACA ATTTCTCCCA GCAGTGCTTG AGAATGGGGA AGGAAAGATA	
9	CAACTICCCC AACCCAAACC COTTIOTGGA GGACGACATG GATAAGAATG AAATCGCCTC	
3	TOTTOCGIAC CGTTACCGCA GTGGAVAAGCT TGGAGATGAT ATTGACCTTA TTGTCCGFTG	
	TGAGCACGAT GGCGTCATGA CTGGAGCCAA CGGGGAAGTG TCCTTCATCA ACATCAAGAC	
55	ACTCAATGAG TOGGATTCCA GOCACTGTAA TGGCGTTGAC TGGCGTCAGA AGCTGGACTC	
	TCAGCGAGGG GCTGTCATTG CCACGGAGCT GAAGAACAAC AGCTACAAGT TGGCCCGGTG	
9	GACCIGCIGT GCTTTGCTGG CTGGAICTGA GTACCTCAAG CTTGGTTATG TGTCTCGGTA	

1560 1620 1680 1740 1800 1860 1920 1980 2040 1140

2100

TTTTACACCA TOCTAAAGAA AAACTTTACA AGGGTGTTTT GGAGTAGAAA AAAGGTTATA AAGTIIGGAAT CITIAAAITIGI AAAAITIAACC AITIGAGIIGIC AAAGITICIAA AAGCAGAACI CATTITICIOC AATGAACATA AGGAAAGACT ACTGIATAGG ITTITITITIT TICICCITIT AATGAAGAA AAGCTITGCT TAAGGGTIGC ATACTITIAT TGGAGTAAAT CTGAATGATC

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CAGAGAAGAG ATTTTTATTA CAAAGAAAA AATTCCAGTG AATTGTGCAG AAATGCTGGT

2160 2220 2280 2340 2395

> (2) INFORMATION FOR SEQ ID NO: 160: 55

CTACTCCTTT GEAGLAAAAC TAGTGCTTAC CAGTTTCCAA TTGTATTTAG CTTCTGGTTG

GAATTTGAAA AAAAAGAAA AAAAGAAAAA GAAAACCTAA ATAAAATAGG TGAAA

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(A) LENGTH: 2120 base pairs (B) TMYE: nucleic acid (C) STRANDELNESS: double

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(i) SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 162:

PCT/US98/04493

30 25 20 5 3 6 357 S TCAGCCTCTC TTACTGTACT CTCCGGGAAT GTTAACCTTT CTATTTTCAG CCTGTGCCAC GAACTETATE CTOTTAGGAT CTTCTGAGET TOTTTCCCTG CTGGGTGGGA CAGAGGACAA CTAGGATCCA GGACTGGGTC AAAGCTGCAT GAAACCAGGC CCTGGCAGCA AACCTGGGAA OCCUPANCIA CATALICACI GACACIGIT CICCICAGAG CICCIGAICT ACCCCACCCC TECETIBEET GIGGGEAGIG GAGAGOETGE TOGGIGTACG CIGEACCIGE CEACIGAGIT CCAGOGIGIT TICCACIAGI CACIACIGIC TICICCITGI AGCIAATCAA TCAATATICI CCCCTCACCA AGGCTGGGAA CAGAGGGGAT GTGGTGAGAG CCAGGTTCCT CTGGCCCTCT GOCAAGGAAC CTTGCTTTTA GCTTCACCAC CAAGGAGAGA GGTTGACATG CCCAGGGCCA CCGCTTCTTT CTTGATCCTC TTTCCTTAAC AGTGACTTGG GCTTGAGTCT CTOTOTAGGO AAGCTGOOTT CCCCATTGGO CCCTOTGGGT CCACAGCAGC GTGGCTGCCC GOCACGAGAT GAGGGGCACC CAGTGCTTCT AGGGCAGGCT GOGTGGTGGT CCCCTAGGTA CAPTICCCTCT GCCCTGAAGC CTGAGTGAAGA CACATGAAGA AAACTGTGTT TCATTTAAAG GTATATATAT ACATATATAT ATTICTTIAA ATTITTIGAGT CITTIGATATO ICTAAAAATC ACCTAGAGTA ANTOGAGAGA CCAMAAGCCT CTGATTTTTA ATTTCCATAA AATGTTAGAA TOGETEGAGG TOGGAGAGAA CETGAACTIC TETTTECCTC TEECTECTEC AACATTACTG AGGAGAAGGG AGGGTCTAGA AGAGGCAGCC CTTCTTTGTC CTCTGGGGTA AATGAGCTTG (1) SEQUENCE CHARACTERISTICS (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 162: (B) TYPE: nucleic acid (C) STRANDEDNESS: doub (D) TOPOLOGY: linear (A) LENGTH: 1003 base pairs STRANDEDNESS: double ACCTOCCOC 1003 960 600 540 480 420 99 360 8 120

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(2) INFORMATION FOR SEQ ID NO: 161:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 900 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear LENGTH: 900 base pairs 5

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GGAGTTTGTC CTTCCACCGA GACTACGAGG GCCTTTGATG CTTAGTGGAA TGTGTGTCTA

2040 1980 1920 1860 1800 1740

ACTIGCICIC TGACATITAG CAGAIGAAAT AAAATATATA TCIGITTAGI CITAAAAAAA

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TGACATETISC ATGAAGCTIGG AGGAGGGCAA ATACETEATE CTEAAGGACE CEAACAAGCA

GTTTGCCAGC CAGATCAACC TGAGCGTGGA GAATGCCTGG GGCATTTTAC GCTGCGTCAT CCACOTOMAA GACTOCTOAC GCOACOTOAT COTAGOCACO CAGCAGITOA AGCOTAATGA

OGICATOCOT GIOTADAGOO TOCOTGATGG CACOTTOAGO TOTGATGAAG ATGAGGAGGA

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50 2 INFORMATION FOR SEQ ID NO: 163:

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GOCCCTCCC GCCGCRGCGA CCTCCAGAGC CTGGGCTACT GCATGCTGAA GTGGCTCTAC

660 600 540 480 420

AGCAGGAGCC CTCACGAGGG GGACCTTGAG TTCATTAGCA TGGACCTGCA CAAGGGATGC

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GAGACCCTOC AGAAGTACCT GAAGGTGGTG ATGGCCCTCA CGTATGAGGA GAAGCCGCCC AAGITIOTIG ATAAGCCGGG GCCCTICGIG GGACCCTGCG GICACTGGAT CAGGCCCTC GGGTTTCTGC CATGGACAAA TTGCCTTCCC AAMAMTGAGG ACATCATGAA GCAAAAACAG

TACOCCATOC TGAGGAACAA CCTAGAAGCT TTGCTGCAGG ATCTGCGTGT GTCTCCATAT

900 840 780 720

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TTCAGTCGGC CCTGGATGTC AGCCCAAAGC ATGTGCTGTG CAGAGAGGTC TGTGCTGCAG

360 30 240 180 120

GITTCGGTGT TCACCAGGAC AAATACAGGT TCTTGGTGTT ACCCAGCCTG GGGAGGAGCC AMBICANCAN GIGGANGANG CIGINCICGA CCCCACIGCI GGCCATCCCI ACCIGCNIGG

AATGTGACAG CTGAAAATAT CTTTGTGGAT CCAGAGGACC AGAGTCAGGT GACTTTGGCA GIOGOCTIGOO GGOTIGOTGAA TIGOOCTIGOAG TITOCTOCATIG AGAATGAGTA TIGTTOATIGA

GOCTATIONOT TOGONTIOCO CTATIOCOCA AGIOGOAAAC ACGIOGOCTA COIGGAAGGO

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SEQUENCE DESCRIPTION: SEQ ID NO: 161:

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CCTCCACCCT CACCTGTRAC TCAGGACCAC AGAAGCAAAA GTTCTCACTC AAACTGGATG GGAAGCTGAA GTCCTTCCAG ACCAGGGACA ACCAGGGCAT TCTCTATGAA GCTCCAACCCA

CCAAGGATGG GCGCTTGTTC AATGAGCAGA ACTTCTTCCA GCGGGCCGCC AAGCCTCTGC

Ε SEQUENCE CHARACTERISTICS: (A) LENGTH: 2196 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 163:

AAGAAGCGGC ACACGGATGT GCAGTTCTAC ACAGAAGTGG GAGAGATAAC CACGGACTTG

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	389		390	
	GOGNAACHTC AGCRIATGCA TGACCGAGAT GACCTCTATG CTGACCAGAT GGAACGAGAA	120	THAITTHAAC TITISTICTIGG ATTISTICCTOT ATTIATICACA OTTICTISTIG AACAGCTITT	1920
•	ATGAGGCACA AACTGAAAAC AGCCTTTAAA AATTTCATTG AGAAAGTAGA GGCTCTAACT	180	CAAGIAITIG GGGAGITIAL CITGCCAICC ICCCCTICIG GITCICIGCA CCCACCIGIC	1980
n	ANGANGANC TECANITIES ASTROCTITI AGGACTICO GATITAACIG AGCICCCIAT	240	CCACTGCAGT TCCTTCCGTG CTCTGTGACT TTAAGAGAAG AAGGGGGAG GGGTCCCCGGA	2040
	AGGAGTACCT GCCTCTTCA GCCCACTAGT AGTGGGCTGG TAAATGCTAC GGAATGGCCA	300	TITIAICITI GITICITITI ICICCTIAGC AGIAGGACIT GAIAITITICA AITITIGGAAG	2100
10	CCTITITISTICS TRACATIVESA TRACSTAGAG CTGATCCACT TITRACCOSST CCAGTITICAC	360	aactaaaaga tgaataaact oggettittit tgetgeftige tittgeaaaa aaaaaaaaa	2160
	CTGAAGAACT TTGATATGGT AATGGTCTAC AAGGACTACA GGAAGAAAGT GACCATGATC	420	arahanah arahanaha arahahara araha	2196
Ā	AACGCCATTC CTGTAGCCTC TCTTGACCCC ATCAAGGAAT GGTTGAATTC CTGCGACCTG	480 15		
3	AAATACACAG AAGGAGTACA GTCCCTCAAC TGGACTAAAA TCATGAAGAC CATTGTTGAT	540	(2) INFORMATION FOR SEQ ID NO: 164:	
	GACCCTGAGG GCTTCTTCGA ACAAGGTGGC TGGTCTTTCC TGGAGCCTGA GGGTGAGGGG	009	(1) SEQUENCE CHARACTERISTICS:	· . · . · . · · · · · · · · · · · ·
20	AGTGATGCTG AAGAAGGGGA TTCAGAGTCT GAAATTGAAG ATGAGACTTT TAATCCTTCA	660 20	(A) LENCTH: 1945 base pairs (B) TYPE: nucleic acid	
	gaagatgact atgaagaga agaggagac agtgatgaag attattcatc agaagcagaa	720		
25	GAGTCAGACT ATTCTAAGGA GTCATTGGGT AGTGAAGAAG AGAGTGGAAA GGATTGGGAT	780 25	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 164:	<del></del>
	GAACTISGAGO AAGAAGCCCG AAAACCGGAC CGAGAAAGTC GTTACGAAGA AGAAGAAGAA	840	GCACAGAGTC GGGGGACGG ACAGGGAGAG GAGGAGAGGG GGTCTGCGCG CGGCCGCTAC	. 09
	CAAAGTCGAA GTATGAGCCG GAAGAGGAAG GCATCTGTGC ACAGTTCGGG CCGTGGCTCT	006	CCAGAAGCCA GCGGACGGCA GCACGGAGTG GGCTGTCCCC GAGCCCAGCC CCGAGCGAGC	120
30	AACCOTOGIT CCAGACACAG CTCTGCACCC CCCAAGAAAA AGAGAAAGTA ACTTCTGAAC	30	CCCCCCCCC CCCCCANGS ACCGCCTYC CAGCCAGCC GACTYCTAGG ASCAGGGGAG	180
	THROCCCTS ACCICCATIC TICCTCCAGC CAACCCTGA AAATTTIACA TGACATAGAA	1020	GOGGGAAAGC AGCTCAAGCC TCACCCACGG CCCTGCCCCC AGCCCCGCCA CTCCCAGGCT	240
35	ACTOTATETY TOCTTFOOTY TICATTICIA GITTICOCAT TICIOTITAT OGCITINGGG	1080 35	CCTCGGGACT CGGGGGGTCC TCCTGGGAGT CTCGGAGGGG ACCGGCTGTG CAGAGGCCAT	300
}	GOCCATITOT GIGGACCAAT CTACTCGGGG AATTCCAGGC CCACCAGGAC ACGTGCCAAT	1140	GAMPHISTIC CIVERCETTICC TICTICCHARCET GCTGGCCCCC ATGGTCCTGG CCAGTGCAGC	360
:	GOCCCCATTC AGATGGCAAG GAAGGAGGTG TTCTTGAAGA CAGGAGGAGG CTCCCGCTGT	1200	TGADAAGGAG AAGGAAATGG ACCCTTTTCA TTATGATTAC CAGACCCTGA GGATTGGGGG	- 420
9	TAATAAATAT TOTTTCATTC TTCTCTTC CTGTCACCTT CTGCCAAGAC ATTGATGGCT	1260 40	ACTIVATORITIC GCTIGROSTICC TOTTICTICATOR CTICGCAGGIC	8
	TCTGACATCT TAITTTOGTOT CTCAAAGCTG TATTTCCAAG ACAGTGGTAC AAGGTGACCC	1320	CANGIGGAGT TTCANTCAGA AGCCCCGGGC CCCAGGAGAT GAGGAAGCCC AGGTGGAGAA	
. 45	TINATIACCE GIATCATGGT TCTTGACCAG CACATTCAAT CCTCCAACCT ACCCTACTGC	1380 45	CCTCATCACC GCCANTGCAA CAGAGCCCCA GAAAGCAGAG AACTGAAGTG CAGCCATCAG	0
	CATGACCTIC CGCACATCTC TAAGITITIAT CITTOCAATA CICAAGGITC TCGGAAATTT	1440	GIGGAAGCCT CIGGAACCTG AGGCGGCTGC TTGAACCTTT GGATGCAAAF GTCGATGCTT	99
	GCTAATGGTT GTGATAAACC ATACAGCTTG AGCCAGTGAG GCAGATTGGG CTGGTGCCTT	1500	ANGAMANCG GCCACTICAG CAACAGCCCT TTCCCCAGGA GAAGCCAAGA ACTTGTGTGT	720
20	COTCTGAGTT TTCCTGCTTT CCTGCCTCGT GCAGATTCTG AGGINTATCT GCTGCCTTGG	1560 50	CCCCCACCT ATCCCCTCTA ACACATTCC TCCACCTGAT GATGCAACTA ACACTTGCCT	780
	AAGACATAAG AAGCAGTGAT ACTCCCTGGC TCGGTTATTT TCTCCATACA ATGCACACAT	1620	CCCCACTOCA GOCTGOGGTC CTGCCCACCT CCCGTGATGT GTGTGTGTGT GTGTGTGTGT	840
55	GSTACANTGA TAGAAGGCAA AATTGCCACT GICTTCTTTT TTTTCTCATA TAICTAAGGA	1680 55	GIGACTOTOT GIOTITICCTA ACTORGOTOT TIGITIGACIAC TIGITITOTOS ATGOTATICI	6
	AGATATATCA GOTTOTOCCT CATGTACCOC TTCTAGTGAA ATGTAGAGGA AGGCTCAAAG	.1740	GITHOTHAGE GAACHGINGA CTCGCTTTCC CAGGCAGGG CTGAGCCACA TGGCCATCTG	096
	GACTCAACAT TTAGATCTÇG AAGOGACAAG TCATGCCTTG GGCCTAGAAT ACCCTGATGA	1800	CICCICCCIG CCCCGIGGC CCICCAICAC CITCIGCICC TAGGAGGCIG CITGITGCCC	1020
9	GAAAAGAGAA GAGGAAGGGA GGCCATATCT ACAACAACAN CCTCTCGGCA CTGCTGCTCC	1860 60		

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35 છ 25 20 5 5 S 50 5 8 8 GCCTAGGGGA TGTCATCAGC ATCCAGCCAT GCCCTGATGT GAAGTACGGC CAGGACGCTT AGAGATTCTT CAGATCCATA CCAAGAACAT GAAGCTGGCA GATGATGTGG CAGCICIACG GCGATTIGGT CGCTTIGACA GGGAGGTAGA TATTGGAATT CCTGATGCTA TEMBELANITT GOOTOGTOMG TOTGMAMOCA ACCITTOGTAM MGCCITTIGMG GMGGCTGMAM CICGAGCIOI AGCAAAIGAG ACIGGAGCCI ICTICITCII GAICAAIGGI CCIGAGAICA TEAAGCCTCC TAGAGGAATC CTGCTTTACG GACCTCCTGG AACAGGAAAG ACCCTGATTG AAGAGICCIT GAATGAAGTA GGGTATGATG ACATTGGTGG TIGITGCICC AGACACAGIG ATCCACIGCG AAGGGGAGCC TATCAAACGA GAGGATGAGG GIGGIGGGAT GCOTGCIGIG GAGTICAAAG IGGIGGAAAC TTAAGCCGTA CITCCIGGAA GCGTATCGAC CCATCCGGAA AGGAGACATT ATGRIGCTIGCC CATTGATGAC ATACTTOTIC TGATGAGAAG ATTCGGATGA ATAGAGITGT TCGGAATAAC CTTGAGCCAG AGTAACCCAT CAGCACTGCG GGAAACCGTG GTAGAGGTGC CACAGGTAAC GACCATTGAT GCCGAGGTCA TGAACTCTCT AGCAGTTACT ATGGATGACT TCCGGTGGGC CTCAGAGGCT GCTCTGCAAG CCATCCGCAA GAAGATGGAT CTCATTGACC TAGAGGATGA ACCTROMACA GIAGOCAATR AGACTICACRG GCATRIROGET GCIRACTIAG CAGCCCIRIES TANAGCAGAG GOCACATOTO ATTOTTATOG CAGCAACCAA CAGACCCAAC AGCATTGACC AAACTOATOG CGAGGTGGAG CGGCGCATTG TATCACAGTT GTTGACCCTC ATGGATGGCC AGATAAAGGA GATGGTGGAA CTGCCCCTGA GACATCCTGC TGAGGCCAAT GTCAGAGAAA TCTTTGACAA GGCCCGCCAA GCTGCCCCCT GTGTGCTATT GTTCTATGGA CCTCCTGGCT GTGGGAAAAC TTTGTTGGCC AAAGCCATTG CTAATGAATG TOCTOTOGAG CACCCAGACA AATTCCTGAA GITTGGCATG ACACCTICCA AGGGAGITCT AGAATOCICC TOCCATCAIC TICATIGATO AGCIAGATGC CATCGCICCC AAAAGAGAGA AAATGTGTTC ATCATTGGCG CTACCAACCG GCCTGACATC ATTGATCCTG CCATCCTCAG GOCTGCTGAC CGAGTCATCA ACCAGATOCT GACAGAAATG GATGGCATGT CCACAAAAAA CCAGGCCAAC TICATCICCA TCAAGGGTCC TGAGCTGCTC ACCATGTGGT TIGGGGAGTC CTOOQAAGAC ATCOOGOOCC TAGAGGATOT CAAACGTGAG CTACAGGAGC TGGTCCAGTA ACCIOCOCOT CITICAICAGO ICAICTACAT COCACTICOT GAIGAGAAGI COOGIGITOC CTITIGATGAG CTOGATTOGA TIGCCAAGGC TCOTOGAGGT AACATTOGAG ATOGTGGTOG ACAGIGGAAG GCATTACIGG TAAICICITIC CCICITIAAG GCAATIGGIG CTGCAGGAAG CAGCTAGCTC AGATICCTAGC CCITATIGCA Tricingico CITICOTOTAC 1800 1740 1680 1620 1500 1440 1380 1320 1260 1200 1140 1080 1020 1980 1920 1860 1560 2100 2040 960 900 600 540 2280 2220 780 720 660

GEAGGECECT TOCGOTTTOT CECCECAGET CECCEMEYEY CYCGRAGAGE CEAGECECEG GATCAGGAGC CAGTOTATAC CGCCCGCCCA CCGCCTTGOT GCCGCTAGAG GAAACGAGAA GOGINGALICE ACGOGINEGG CAGENGTINGT TINGAGINGTIT GETTGENGTIG CONCENTENCES ACACAGIGIT GCIGAAAGGA AAGAAGAGAC GAGAAGCIGI TIGCAICGIC CITICIGAIG E Œ SEQUENCE CHARACTERISTICS: GCGCGCCATG GCYTCTGGAG CCGATTCAAA AGGTGATGAC CTATCAACAG GGTGTCCTTG TCCCAGCCCA AGATGGATGA ATTGCAGTTG TTCCGAGGTG ACAGAAGAAC COTCCCAATC GOTTAATTOT TOATGAAGCC ATCAATGAGG COCYTOCCAC CSCTCOTAGC COTTACCCGC GOOCCGCCAC AGCCGCCGGC SEQUENCE DESCRIPTION: SEQ ID NO: 165: (A) LENGTH: 2933 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear

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480 420 360 300 240 180 120 3

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(2) INFORMATION FOR SEQ ID NO: 165:

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GGTCATAACG AGAGTGGGAA CTCAACCCAG ATCCCGGCCC TCCTGTCCTC TGTGTTCCCG ACCTICCCIG CITCIGAGAC TICAATCIAC AGCCCAGCIC ATCCAGATGC AGACTACAGI GIGICCCCIG CATATCTICT CAGCAATAAC ICCAIGGGCT CIGGGACCCT ACCCCTICCA CCCCTTCACA GAGCGCCCGG GGATTCCAGG CCCAGGGCTT CTACTCTGCC CCTGGGGAAT ANGCOCTROC GROGITAANT TOTROCCAGG GGOTROCACG AGGAGROCCOC ARCTGOCCOG TTAGCAACTG GAGATACAAA GCAAGGAGCT GGTGAGCCCA GCGTTGACGT CAGGCAGGCT TICACICCII TAACAAAAAC CITOCIICCI TATCCCACCI GATCCCAGIC TGAAGGICIC CAGINGINIT GOGACCIOGG AAGOTTIGCA GCACITIGIC ATCATICITC AIGGACINCI CGAGOTOGOT TOGAGACTCA GCAGGCTCCG TOCAGCCCTT OOGAACAGTO AGAGOTTGAA TGGGCAGCAG AGGCAACTCC CGCATCCTTT GCTCTGCCTG TCRGTGGTCA GAGCGGTGAG ACACCGGGAT GGATGGAGGG AGAGCAGAGG CCTTTGCTTC TCTGCCTACG TCCCCTTAGA COCTOCAATT GGGTCTCTGG CAGGCAATAG TTGAAGGACT CCTGTTCCGT TGGGGCCAGC COGRARACCAA CCARACCOTO COCTOTGACC CATTGCTOTT CTCTOTATCG TGATCTATCC 1920 1500 1945 1860 1800 1740 1680 1620 1560 1440 1380 1320 1260 1200 1140

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GAGACCAGCC CCCTCCCCTO ATTTAGOGAT OCOTAGGOTA AGAGCACOGG CAGTGGTCTT

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CATECTEANG OCTANCETOC GEANGTECCE AGTTOCCANG GATOTOGACT TOGAGTTCCT

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	393		394
			AAGATITIT AAAATAGGG CTGTGTTTAA AAAAAAAAC AAACARGAA AAGCAGCAGT
	GOCTANANTS ACTANTOSCT TCTCTGGAGC TGACCTGACA GAGATTTGCC AGGGTGCTTG	2340	GATTATAGAG AGSTCACACT CTAAGTGGGG TCGCGGCGTG GCCACGCTTC ACGSTCACGC
٧٠	CARGCTGGCC ATCCGTGAAT CCATCGAGAG TGAGATTAGG CGAGAACGAG AGAGGCAGAC	2400	TOSTOCETICC TECASTIGACE TOTITIACATE STCACACOTO TOTISTATCAC CAGTIGASTICA
	AAACCCATCA GCCATGGAGG TAGAAGAGA TGATCCAGTG CCTGAGATCC GTCGAGATCA	2460	ACTIGCTIGIC ATTCCTCCCG TGGCAGTTTG TGTAGACAAT CTTACTGAGC AAAAGGCAAT
:	CTTTGAAGAA GCCATGGGGT TTGGGGGCG TTCTGTCAGT GACAATGACA TTGGGAAGTA		GAAAGICIT GGITCCCACA CIGCGAIAIA TIGGAAITIT CACCICAGIT TAIGAAGITI
0	TOAGAITITT OCCCAGACCC TTOAGGAGAG TCGGGGCTTT GGCAGCTTCA GATTCCCTTC	2580 10	ATTICGAAAF CCATAGTCAT CTAAGAATGA ATACCTGTCT GCCATGTATT TCAATCTTAG
	AGGAAGCAG GOTGGAGCTG GCCCCAGTCA GGGAAGTGGA GGGGGCACAG GTGGCAGTGT	2640	TGAGCCAAAA TIGITIIGITT GITACTACAG AATAGAGATG ACTGITITIT GCCACAGCCC
15	ATACACAGAA GACAATGATG ATGACCTGTA TGGCTAAGTG GTGGTGGCCA GCGTGCAGTG	2700	TATIOGRAFIT GCAATCTOTO ATTGCCTTOT AAAAAGAGA GTGCATATGG CACTGCATTA
	AGCTGGCCTG CCTGGACCTT GTTCCCTGGG GGTGGGGGCG CTTCCCCAGG AGAGGGACCA	2760	AACGTGTGGT GTTTCTAGTC AATGATATTG GTGAGCACAA TOTATTCATT TAATGGCATA
	GOOGTOCOCC CACAGOCTOC TCCATTCTCC AGTCTGAACA GITCAGCTAC AGTCTGACTC	2820	GACCATACCA GACCTAATTT GCAAGTATTG GGTCTTAAAC TTCAAGTGCA ATGTATAA
70	TOGACAGOGG GITTCTOTTG CAAAATACA AACAAAAGC GATAAAATAA AAGGGATTTT	2880 20	AAACCAARCT GAGCCTICPA TCTCTTAAAT ATTTATTTTT TTTAACGTGT GAGATGTTCG
	CATTIGGTAA AAAAAAAAA AAAAAAAAT CCGGGGGGGG GCCCGAACCA TIT	2933	AGAGAAGTI CICCAITCAI TICAGIGCIG CCIGGAGGAA ACTOGGCAAI GAITICITIC
25		25	AGTISTISMAG TYCCTTTCGF GTIACACCCT CCACTGAACC CYCAACCTTC GAAATACTCC
	(2) INFORMATION FOR SEQ ID NO: 166:		AGITITICIES GITICOICAT ITITACITAL AAAITIACCI TITIGIATIT ISCAAITIAC
ć	(1) SEQUENCE CHARACTERISTICS:	QF.	AICTOTITICS TITICITITAA AITCICIGAA AGIGOCTICA TIAAAAGACT CCITITIAAAT
3		or .	GGAAGCCACC AGTCAGCAGA ATGGAAGCTT AGAGGAACTT GCCTGTGAGC GCTGGTCTTT
	(C) STRANDEDKESS; GOUDLE (D) TOPOLOGY: Linear		CICITICGIT INCICATORA ACGRICITIG CICCOGNITY TICCTITICIT INCAGGGAAA
35	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 166:	35	IGICITGGAG TAAAITITAA GITCCTGGAG TIAAITITGIT TIACAGGAAF TITGITITIT
	TOGGAGAGCC GACGGAGGNG COCCTCTCGG CCAGGAAGCG CCTCTTGGAC GCGTGTNACC	09	AAAAAATAG GATCATTCTG AACTTTGGAA TGACCCCCTT ATATATTTTC TGAAAATGAA
Ş	GATOCCCAGA AGTOGCCTTG GOCTOGGGAT CACCATAGCT TITICTAGCTA CGCTGATCAC	120	AACAGTTACA TGAAAAAAT TTCCAATGAA GATGTCAGCA TTTTATGAAA AACCAGAAGT
₽	GCAGTITCTC GIGTATAATG GTGTCTATCA GTATACATCC CCAGATITCC TCTATATTCG	180	TATTAGATGA AAGCAGGGAG TGAATCTTTA AAACAGACTT GATCAGGCAC ACACAATAAG
	TICTIGGCIC CCTIGINIAI ITITICICAGO AGGCOTCACO GIGGGGAACA IAGGACGACA	240	TCTTTCTCTC CGAAACCGGA AGTAAATCTA TATCTGTTAG AAATAATGTA GCCAAAAGAA
45	OTTAGCTATO COTOTTCCTG ANABOCCCCA TAGTGATTGA GTCTTCANAA CCACCGATTC	300 45	tgiaaaitig aggattitit toccaatagt tiatagaaaa tatatgaacc aaagtgaitt
	TCACAGCAAG GAACATITITG GAAGAAAATC TCACTOTGGA TTATCACAAA CALTAICTTT	360	GACITICIAA AARICIAAAA TACIRICAAC AAAAITIGCA CICIRCCAGA ITIGAACAIC
Ş	TITCITAAGT AAICIATITA GAICGGGCTG ACTGIACAAA TGACTCCTGG AAAAAACTCT	420 50	TAGICAGGIT CACATICATA CTAAGITITIC AACATIGIGT TCTTTTIGCA TICATITITI
3	TCACCTMOTC TAGAATAGGG AGGTGGAGAA TGATGACTTA CCCTGAAGTC TTCCCTTGAC	480	actititatia aagoticaaa acc
	TOCCOGCACT GGCGCCTGTC TGTGCCCTGG AGCATTCTGC CCAGGCTACG TGGGTTCAGG	540	
55	CAGGIGGCAG CITCCCAAGI AITCGAITIC AITCAIGIGA TIAAAACAAG TIGCCAIRIT	600 55	(2) TABORAMAPTAN FOR SEO TO AND. 167.
	TCAAAGCCTT GAACTAAGAC TCAATTACCA ACCGGCAGTT TTGTGTCAGT GCCCAAAGGA	099	(1) CENTIDALTO PLAN APPROPRIATO.
09	остасства тостоства сласатва статестота атассатка таттатеса	720 60	(A) LENGTH: 1816 base pairs (B) TYPE: nucleic acid

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GTAGAAATGA TOCTTCTGCT CAGGAATGGC CCACAAATCT GTAATTTGAA ATTTAGCAGG 1680

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INFORMATION FOR SEQ ID NO: 169:

(1) SEQUENCE CHARACTERISTICS:

(B) TYPE: nucleic acid (A) LENGTH: 902 base pairs 50

GTTAACCATT ATAGAAAAGC AAAGCTTGAG TTTCCTAAAT GTAAGCTTTT AAAGTAATGJ

900 840 780 720 660 600 540 480 420 360 300 240 180 120 S

TCTTAAGAAA AAAGOGAGAA ATATTAATCA GAAAGTTGAT TCTTATGATA ATATGGAAAA

ACATTAAAAA AAACCATTAT TICACIGICA TITAAAGATA AIGIG

45

GGTAATTCTG GCATGTCCTC AAAAATGACT CATGACTGTG GATATGAAGA ACTATTGACT

CIGATOCIGA TITIOCACICI OCIOGAATIC TOCCTAGCIG IGCICACIOC TOTOCIGOGG

ATACCACGGA CTOCTATACA GCCAAAGCCA GTCTGGCTGG AWCTCTCTCT

TOGANACAGO CITACICIGA CITOCCIGGG AGIGIACITI TOCTOCCICA CAGITACATI

6

GATTCACTT

CAGTOTGAGT TOGACAAAAA TAATATACCA ACAAGAAGTT ATGTTTCTTA CTTTTATCAT

35

accendance

30

CCATTCATAG GACCCTITIT TITTATCATC TCTGGCTCTC TATCAATCGC CACAGAGAAA

TCTGCTTCCT TCTCTCCAAA TTTTACCCAA GTGACTTCTA CACTGTTGAA CTCTGCTTAC ATTOGOACTA TCCAGATCTT GTOTOGCATG ATOGTATTGA GCTTOGGGAT CATTTTTGGCA

AGGITRACCA AGCITITIGGI GCATAGCAGC CIGGITIGGAA GCAITCIGAG IGCICIGICI

GTTTCATTAT CCTOTCTOTC AMACAGGCCA CCTTAMATCC TGCCTCACTG

25

GAACCCACCA ACCAGGGGCA GGATAGCCTG AAGAAACATC TACACGCAGA AATCAAAGTT

CAGCTOGAAC CAACGGGCAC AGTTGGCAAC ACCATCAACT TCTCCCAAGC AGAGAAAACCC

AGAAACCOTT GATOOGACTG AGAAACCAGA GTTAAAAACCT CTTTGGAGCT TCTGAGGACT

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SEQUENCE DESCRIPTION: SEQ ID NO: 168

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 945 base pairs
(B) TYPE: nucleic acid
(C) STRANDEINESS: double
(D) TOPOLOGY: linear

INFORMATION FOR SEQ ID NO: 168:

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GGTACCCUAT TCGCCG

AATTATGTGC TGAAAAAAAA AAAAAAAAAA AMMMRARASK RRWWACTCGA GGGGGGGCCCC AAATGACCIT TAATGACACI ACATTITICAG GAACIGAAAT CATTAAAAIT TIATITIGAAT

1816 1800 1740

8

CYSCTTCTCA CCTCCTGCCA TGATACTGTC AGTTACCTTA GTTAACAAGC TGAATATTTA 1620

GCCCGTTACC AAKTCGCCCT AIWGTGADTB GTATIMITAT TITACTAATA TCTGTAGCTA KGCTIKGGIT ATKGTTTITY TCCCTTYTCT WAGCTATRAG CTGATCATKG 1560 1500

55

50 TCATAGAGCT TTTAAAATGG TTTCATTGGA TATAGGCCTT AAGAAATCAC TATAAAATGC 1440

TANAACTGAA GATCATGAAG AAGCAGGGCC TCTACCTACA AAAGTGAATC TTGCTCATTG TURARTITIAA GCATITITICI TITIARAAGAC AAGIGIRATA GACATOTARA ATTOCACIOO 1320 1260

45

CACGCTGTTG GACGCAGTAT AGTITICCCTC TGAGAAGCTG AGTATCTATG GTGACTTGGA CTTTATGAAT GAACAAAAGC TAAACAGATA TCCAGCTTCT TCTCTTGTGG TTGTTAGATC GICCICICGG IGAIGGIAIT GCITIGGAIT IGIIGIGCAA CIIGIIGCIA 1140 1080 1200

8

AGATOGAGAA AGTGATOOCT TITTAAGATO CCTCTCTCTT AACTCTGGOT GGATTTTAAC 1020

AAGCAAAATG TCCNTCAKMT CGSMAATGAG AAATTCACAA GCGCACAGGA ATTTTCTTGA TOCCAGOTAC GOACCACATT TOGAGOCCAGG AGCCCTACCA AATTTORGRG RAMONTOTOT 960

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GACTITITAT CTICAAGCCG ATGACGGAAA AATAGITATA TICCRGTCTA AGCCCAGRAA 900 840

GOTGAGGTCA TICTGGAGTO ACATGATOGA CICCGCACAG AGCITCATAA CCICTICATO 780

30

720

GAGACAAGAA CAACITATGT CCCTGATGCC AAAAATGCAC CTACTCTTTC CTCTAACTCT ATCTGATGAG CAATATGCTT GCCATCTTGG KTGCCAGAAT CAGCTGCCAT TCGCTGAACT 66

AATTGACTTA AATCGAACTA AATTGGAATG TGAATCTGCA TGTACAGAAG CATATTCCCA 600

25

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GTTGTACGCA TOTCAGAGAG GTTGCAGGCT GTTTTCAATT TGTCAGTTTG TGGATGATGG

540

8

GICTIGCCAC COGOCCIGIC AGTIGACCIA CCCCITGCAC ACCIACCCIA AGGAAGAAGA GOCCOGAGOT TCGGGGACCG CTTCGGCTGA AGCATTTGAC TCGGTCTTGG GTGATACGGC 420

GAGCCTCTGG GTGAGGACCC AACTGGGGCT CCCGCCGCTG CTGCTGCTGA CCATGGCCTT 360

GAAAACAGCA ACAAGCTGAG CTGCTGTGAC AGAGGGGAAC AAGATGGCGG CGCCGAAGGG GENACUTICO GTUGGOGGA GAAGGGGCT GGCCCCAGGA GGAGGAGGAA ACCCTTCCCA 8 240

COCAGGAGOC GOCAGITITET GOCGGGTGAG GGCGGAGCTG AAGTGACAGC GGAGGCGGAA 180

5

TICGGGICGA CCCACGCGIC CGGCCAGCCI AGGAGAGAA GIICGIAGIC CCAGAGGIGA 120 GOTGGONAGO TITINAAITITO COCTTACMOG GGCOCTNIAA GGGGAAAACCI TOCCGGAATT SEQUENCE DESCRIPTION: SEQ ID NO: 167:

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(C) STRANDEDNESS: double (D) TOPOLOGY: linear

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WO 98/39448

PCT/US98/04493

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WO 98/39448	4	PCT/US98/04493	WO 98/39448	PCT/US98/04493
	397		398	
(C) STRAND	(C) STRANDEDNESS: double (D) TOPOLOGY: linear		TCCTCCTTTG TGGTAGCCTC TGGCCATCTG GGACCTCAAT CCCCAGCTTT CCCACTTTCA	360
אמת מטותחוטמט (יָרי)	SALL CONTENTS DECORPORATED CON TO NO. 150.		GCAGTCCTTT GCTCTCTTTG CTTCTACCTC AAATAGCCCC AGGAGTGGC TTTAGTCTCC	420
(XI) SELUENCE DES	CALFILLAN: SEQ 1D NO: 109:	ν,	ARTATIGEAGC ATTICAAGCT TCTCCTGGGG GATGGGGATT GGGATGGGCA GAATCTGTTT	480
GOCALAGECCA CAGGAAGGAT GAG	KACAGAKCCA CAKGAAKKAT GAKGAAKACC AKKCICIKKK KACIKCIKI GATIKCICIT	00	TOGNICICOS GOTINITICO ACTOGOTOTA AAAGCAGAGO TOGOCOTTIC COTOTOTAT	540
	GICILMAAR TUCGAICTIC AACTAANTI'A ACTIGAIGAAA AGTATGAACT GAAAGAIGGGG		CCCTCAGGGT GGGTAAGAAG GACTGTATCT ACACCTGTTC TTCCCTACCT TCTCTTTTGT	. 009
10 cagaccerss argigadang mga	CAGACCCTOS ATGTGAAATG TGACTACACG CTAGAGAAGT TTGCCAGCAG CCAGAAAGCT		TAGGGAGGC TCATTCTAAG TTCCTCAAGA GAGTCCTTGG CTTAAAGCTG TAGCAAGGGT	099
TGGCAGATAA TAAGGGACGG AGA	TOSCHOATAN TANGGAIGGG AGNANTOCCC ANGACCTIGG CATGCACAGA GAGCCCTTCA	240	GIOCTAGGIG CAGGATITICG ACCAAAACCG TCGAGTAGGC ATGATACTICG TAITCGAGTICG	720
AAGAATTCCC ATCCAGTCCA AGT  15 TTACTGGGG TCCGAATGGT CAA	AAGANTICCC ATCCAGTICCA AGTGGGGAGG ATCATACTAG AAGACTACCA TGATCATGGT. PPACPICCCG TCCGAARGST CAACCTICAA GTGGAAGATF CTGGAACTISTA TCAGTRITISTS	300	GCCTGCAAAA TCAGACAGAA ATGGCTTGAG AAGCCGCAGG GGAGCATGCC TGTCTCTCAG	780
ATCTACCAGC CTCCCAAGGA GCC	ATCHACAGE CICCAAGA GCCICAAAA CISTICAACA GCAICCCTT GSDGSTGAC	420	TGHTHGHGTA TGGGHGGGHC CTCCCTHGCT TGGANANTGH GANTTGAAGG GGTTHTGAAC	840
20 saccomme cacesacon nee	AACSCEPTITO CACCOACTO TOCOTOCAAD CACAATITOTA COCCAANTO COADAACATO	480 20	ANATHGEATG CCTAOTTGAG GATGTTCCCA AAGTTTTGTC CAATCTTATC ATTAGTAGAT	006
	CETECTACEA CALTANDOC CITUTOCOCCA CICTATACEA GOOCCAGAAC TOTGACCCAA		TITIATAAGCC ACAGAGACAA ACCAGAAACG GAATAATGTT ACTITIGGATG CTTTATTTTT	096
	SCHOOLOGE ACTENACTIC CONTESCUES ACTECCTORACT CTOLANTICAA CCTTACAAAT		ttottotagg totgacitig tacatgcaga agaatgctat atgctgcaca ttttgccttt	1020
25 GTGACAGATA TCATCAGGGT TCO	STRACHANTA TCATCAGGGT TCC003T07TC AACAITTOTCA TTCTCCTGGC TGGTGANTC	25	AAAGHCTTAC GACTTTCCCC ATTTTAGTCT AATGGGAAGA TACAGATGTG CAAGTCTGCT	1080
CTGAGTAAGA GCCTGGTCTT CTC	CTGAGTAAGA COCTGGTCTT CTCTGTCTG TTTGCTGTCA CGCTGAGGTC ATTTGTACC	720	TTTTIGTTT TIGTATAT TTTTTTTT TIGCICIGIG TAIGGACAT TTTCAGACAT	1140
30 TAGGCCCACG AACCCACGAG AATA	TAGGCCCACG AACCCACGAG AATGTCCTCT GACTTCCAAC CACATCCATC TGGCAGTTGT	780 30		1200
	GCCAAGGGAG GAGGGAGGGG GTAAAAGGCA GGGAGTTAAT AACATGAATT AAATCTGTAA		TAICAGCIAI GGICAACCIG GTITCAICIG TAICICICIC TITICACCIG TAITGITIAI	1260
מפר ממפתמתמתמת מדייניםיייתייי	HOROCOGOTA RARARARARA RARARARARI CCAMPOTANCO INTITIONIZATIV CRITCRIZATIV	CCC	TGAAAATCCA AGACACTATG CCAATGCAAC COTGACTACT TTOGGAGAIT GOTAGTCTCT	1320
35 TT		35	ITTGATGOTG ATAOTGATISG GOTGCACTAT CATAATCACA TCAGOTCTSC TTTTTGCTTT	1380
<b>.</b>		•	taangitaac taangaagit ccagagangg gccttagaaa tgngittiaa gaattaacaa	1440
ç		40	GGAGTETEAA AAAGAAATGA GAGGGATGET TECTTTEECE TTGCATETAE AAAGAAGAG	1500
+0 (2) INFORMATION FOR SEQ ID NO: 170:	D NO: 170:		AGAGACTOTT CTGTTGTAAA ACTCTTTCAA AAATTCTGAT ATGGTAAGGT ACTTGAGACC	1560
	E CHARACTERISTICS:		CITCACCAGA ANGICAANCY TITITICIGI GIAACATGGA AACTIGIGIG ACCATIAGCA	1620
(A) LENGTH: 45. (B) TYPE: n	LENGTH: A003 DASH PALES TYPE: Mucleic ecid	45	TIGITATCAG CITGIACTGG TCTCATAACT CTGGTTTTGG AAGAATAATT TGGAAATTGT	1680
	Sironucanos; double TOPOLOGY: linear		TIGCTOTICTIC TOTGLADATA ACCICCCCAA AATAATTAGF AACTOGTIGF TCTACTTIGGF	1740
(xi) SEQUENCE DESC	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 170:	05	AATTIGACAC CCTOTTAATA AGGCAATTAT TICTIGIGTIC TIAAACAGTA TAAATAGTIG	1800
	AGAAAACAAC TGAAAAACCA CATTTTTCTA CATACAGCTG GGGAGGTAGC TGAGAACTTG	09	taagitigca igcaigaigg aaaaataaa accigiaici cigitaaaa aaaaaaaaa	1860
GCACTGGGCA CACATACTAG GTT	GCACTGCGCA CACATACTAG GTTGAAAGAG AGTTGAGGAA ACCAGAAGGC CAAGTGGATC	120	AAAAAAAA AAAAAAAAA AAA	1883
55 recreaceae cocreaect orc	TGCTGGCAAA CCCTGAACCT GTCTCCTGCG CTTGCTCTAC AGTTCTGAAG TTGAAAATCC	180 55		
TITICATGCC TAGGATCTGC TTG	TTTTCATGCC TAGCATCTGC TTGAGTTATA AACCCCAAGG CAGCCATGTC ATAGACTAGT	240	13) THEOMILIAMENT DAD GEO IN 171.	
gmacrem grrmaadm ner 60	orreacter orresoler torresole cerectogo ecologice rectoriors	300		

WO 98/39448

LENGTH: 2100 base pairs

9099 TYPE: nucleic acid STRANDEDNESS: double

Œ SEQUENCE DESCRIPTION: SEQ ID NO: 171:

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55 50 દ 6 35 છ 25 20 5 5 GATACAGAGA TITITITITT CCTTGGAAAC AAATGGACTG GGAAGAAACA CAGCATGGCT TITICIGITY TIACCITGIC CAATIICCAC ACTAGICATI TITITIATIT TITAGAGGAT CICTIGGIGT CCICTIGGIC ATAMAGITOT CICCIACCIA ACCCAGITIT ACCAMANGGA GAAGGGAGGA CATGATTCCA AAAAAGATCG TICTCAATGT GICGTCTGAC TCAACCAGCT TACATATOTA TOTACACACG GTACCCAGAG TOOTACTOTG GCAGCCTTCA CAGATITITAG CGCTGGAAAA TGAGTTCAAA AATTTCAGTG TAATGTCATA AGGATGTTGG CICITIOGCA AMAGGGIGAT ACTITICACT AMAMAIGCCI ACICITCCIG TIGAIGITICC CATAATTICA AGAGIGIGAC CACCCIGCIC TAGICAICAT CATIGGATGA AICCAGITGA CTTTTCCATA TAAGGAGCCC CATTACATAA GCTACGGGTG AGGTTGGAAC AGCTATGTTT TTGAATATGC CTCTGTTTGG GCAAAGCAAG ATACCTCCAC TTAACCTTTA TCCAAGGAAG GCCAGATTAC ACTIGCCAAG TOSTICCCTT TOCTICIAAG TOAGTIGGCT COATATICAC ATACTOTTGA TOGAGCAGCA GCATAGCCTA GAGTGATGCA TTCTTACCCA GAGGTGGCAA CCTITITOTCT CTCAGGCTAA CATGAGAGAA AATAGAAAAG TCTTGGCTGT GGGGATTGGA GIGITICIATA AACIAAGCAI COGIOGGITT AGAGIGITAA AGIGICAGCA CAITCCIICI AAAAAGATGT TIAAGAGAAT TAATAGAGCC GTAGTCTGTA TIAGGATGTG TGTCATATGT TACTITIAGA TITACIOCCI TCAAAAAGIG CCIATICIGA GCAACATAAA AGTAAAAGGG GACAAACTAT GGAAGATGGA CTCCATGCCA TTOCAGTCAG CCACCATTCT TAGGAGAGGG TCCATGTAAA TAGGACGAGG TAGACAGTGC ATGATTGTAG GAGAAGGGTT AGTAGTGTAT GITATITIGAT GGCTTITIGIT TCCATAGTTC CATCACTGAC AAAACTGTCA AGCTICAGGG GCCAAATIGIC CTTIGCCAGAT CCTTIAGAGCA TTACTTTIGAC TCCTAAAAAT GTTTCCTGTA CTCTGCTTCA AGGGAATGTA AGYTTTATGG CATTGAAAACA TTTAGGAAAA TOCOTTOCTO ATACAGOTTO TITTATATITA TATOCTACIO GATOGTAGOA TATIGOTAAG ATCAGAAAGA GTAGGTGCTG AGATAAGGNA CCTCTTCAAG CGGCTACCGT GRAASGGGCT GCAAACACAT ACTITIGCCAA AIGNAAGAAA GICACICACI CGITATICCI 1440 1260 1200 1080 1020 1320 720 660 600 540 480 420 360 300 120

> 2 0 TATAAGCAAG TGATTTAGGT ATTTTCTTTT GTGTTTATGC ATTATCTGAC TATATTAAAA ATAAGAATGA ATATGGAAAT TATATTTCTT TTTTCTGTAA AAGAGTTGCA ACTACTITAT CACTGATOTA TCCAAAATAG CACACATAGT TCAGTATGAA AATAAGAGAA TAAAATCTGT GOTTICTCTT TITTICTTGTA TICTTAGCAA ATTGCATTTA TICACTACAT TACAAACCAT CCIGITITIC TATTIACCIT CIATCAGITI ICICIACCAA ITAIGITITI ICAAIGCICI AAGAATATOT CTATTTTCAT ATGTGTGATA CTGACAGAGC CATGGTATTC CTAAAATATA GATGAGCTGC TAGTCTGAAT AACATTCCCT GACTTAGGGA AAGGCACACA AAAACATATA 1740 1680 2040 1980 1920 1860 1800

E SEQUENCE CHARACTERISTICS: (A) LENGTH: 1930 base pairs (B) TYPE: nucleic acid (D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO:

172:

8 ઝ 30 ACAGITITIG GOTOCTOGTO GTOGGGGGGG GCGGCATCGG CTGCGAGCTC CTCAAGAATC TCGTGCTCAC COCCATOGCA CTOTCOCOGO GOCTOCCCCO GUAGCTOCCT GAGOCGOTOG CCGGGGGCCG COTTIGANTS TOSTICCOGG TICCHGATTING CASCOCCTICC GCCCCGGCTC GTGGTTNTCC COGTTTCTCC Ĕ TTTCAAAAGA AACATGTTGG AAGATCAAAG GCACAGGTTG CCAAGGAAAG CACATOGACO TGATTGATOT GGATACTATT GATGTAAGCA ACCTCAACAG SEQUENCE DESCRIPTION: SEQ ID NO: 172

ACCIDECEGA AACCAITOTTA ATAGAATISTO CETEGEAGET GATOTTECTIC TTATTIGAAAG AACGGAAGCC TEMPTOTICAT COTAMBECEGA CECHGAGAAC CITTECTOGO TOTACAATTO GIAACACACE TGACTATAAT GIGGAATTIT TCCGACAGIT TATACIOGIT AIGAAIGCIT TAGATAACAG TAAAGATGAC ATCAGGTATC TGTTGACAAT GGACAAACTA TGGCGGAAAA GGAAACCTCC AGAAGATGOT GATCAAGAAG TATCTOOTGA CAGAGOTGAC COTGAAGOTG COTGGGAACO TICAGAACCI ATACATICCA TCGITIGGGC AAAGIACTIG TICAACCAGT IGITIGGGG PACTAAGGAA TGGGCTAAAT CAACTGGATA TGATCCAGTT AAACTTTTTA CCAAGCTTTT GAAGCCAGAG CTAGAGCATC TAATGAAGAT GGTGACATTA AACGTATTTC GOSTATETTO GACAMSTANC TACTATEMAA AMSSSTSTGA CEGAGTSTTA TITIACCCGA AAGCTAATAT CGITGCCTAC CATGACAGCA TCATGAACCC 300 240 180 120 900 780 660 600 540 480 420 360 840 720 60

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TANATOTOTO TACTINAGAGO AGCACTTOOT ACTAGOTAAG CACAATCATA GCCCCACCGT

ATGGTGTTTA CAAGTGGATA GATAAGGCGG AGATGGTGAG AAGCCGGGTT TTCTCTATGC

1560

1500

1620

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TIGCICIONG TITCANICIG AIGATIAIGA CCAIGGAAGA TAGICITAIG TAAAGGITA

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	WO 98/39448	PCT/US98/04493	WO 98/39448
			402
	AGTICCOTTO CACTOCOCTO AAGTACAAAG TCAAGGAGAA GAAACGAATO CATCAGATCA	096	TOTGGCCCAA AAGAIGGCTG AGCCAGAGAA GGCCCAAGC CTCAGAANT TOTAGAANT
	ACAGANTGAA CCCCAGTTAG GCCTGAAAGA CCAGCAGGTT CTAGATGTAA AGAGCTATGC	1020	
S	ACOTETITIT TCAAAGAGA TCGAGACTIT GAGAGTICAT TTAGCAGAAA AGGGGAITGG	1080	S GACACTOCTO PTERCENCE POSTANTEM CONCENSES
	AGCTGAGCTC ATATGGGATA AGGATGACCC ATCTGCAATG GATTTTGTCA CCTCTGCTGC	1140	ACTICATORIA MANAGERIA ACCORDANIA ACCORDANIA ACTICATORIA CALCINA ACTICATORIA AC
2	AAACCTCAGG ATGCATATTT TCAGTATGAA TATGAAGAT AGATTTGATA TCAAATCAAT	1200 . 100	COCCONCIO CALTTERAN CANTONINA CONTRACTOR ACANTOGOGO
?	GGCAGGGAAC ATTATTCCTG CTATTGCTAC TACTAATGCA GTAATTGCTG GGTTGATAGT	1260	
	attogaaga ttgaagattt tatcagaaa aatagacag tocagacaa tttttttgaa	1320	
15	TAAACAACCA AACCCAAGAA AGAAGCTTCT TOTOCCTTOT GCACTGGATC CTCCCAACCC	1380	
	CANTIGITAT GTATGTCCCA GCAAGCCAGA GGTGACTGTG CGGCTGAATG TCCATAAAGT	1440	GCCTISTOTS GCCOTCAACT GCCTGTCAAG CTCACCGGGT ACCCCAGGCC ACAACTAAT
20	_	1500 20	_
	TOTCCAAATT GAAGATGGGA AAGGAACAAT CCTAATATCT TCCGAAGAGG GAGAGAGGA	1560	TOTITIANC TCCTITIGGT ACCTINATIT CACTORCCIC GCAGAGAAM REALANCE
	ACCTARINAT CACANGAAGT TGTCAGAATT TGGAATTAGA ANTGGCAGCC GOCTTCAAGC	1620	Ī
22	AGATGACTIC CICCAGGACT ATACTTTAIT GAICAGAIC CITCATAGIG AAGACCTAGG	25	
	AAAGGACGIT GAAITIGAAG TIGITGGIGA TGCCCCGGAA AAAGIGGGGG CCAAACAAGC	1740	GITOTOCOAC CCITICITIA PARTICIAN ACCIONACION CONTINUEDE PARTICIANO DE CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDE PARTICIANA CONTINUEDA PARTICIANA PARTICIA
20	TGAAGATGCT GCCAAAAGCA TAACCAATGG GCAGTGATGA TGGGAGCTTC AGCCCTCCAC		
2	CTYCACAGCT TCAAGGAGGC AAGATGGACG TYTCYCATAG TTGATYCGGR TGAAGARGET		
	TOTOCA APPARA POTOCOCCA CONTRACTOR CONTRACTO	0000	GACATOTTOC TOCTAATOCT GAGGCTOGTA GCAGAATOCA CATTOGAAGC TOCCACOCCA
35	TITCH THE TRACE OF THE TRANSPORT OF THE TRACE OF THE TRAC	1920	TATTGITCIT CAAAGTGGAG GTCTCCCCTG ATCCAGACAA GTGGGAGAGC CCGTGGGGGC
3	TTAGANTIG	1930	AGGGGACCTG GAGCTGCCAG CACCAAGCGT GATTCCTGCT GCCTGTATTC TCTATTCCAA
			TARAGCAGAG TITIGACACCG TCAAAAAAA AAAAAAAAA AAAAAAAAA ATTACTGGGG
40	(2) INFORMATION FOR SEQ ID NO: 173;	40	CCTCAAGGG
45.	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 1509 base pairs  (B) TYPE: nucleic acid  (C) STRANDELNESS: double  (D) TOPOLOGY: linear	45	(2) INFORMATION FOR SEQ ID NO: 174:
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 173;		(1) SEQUENCE CHARACTERISTICS: (A) LENTH: 177 base pairs (A) LENTH: 177 base pairs
20	GACCCTGGCC TCTGGGCTGA GGCTTGCTAG GGACTCGGGG TGGCTCTAAG GGGCAGGGAT	99	(a) iffer intercording (c) STRANDEDES; double (D) TOPOLARY : ileas
	AGGGCTGGGG AGCGCCGGCC TGTGGCCCTG ACCAGCCCT TCTCGTGCRG GTTCCACCCC	120	
55	GATOCAGOTIC STCACGTGCT TGACGCGGA CAGCTACCTG ACCCACTGCT TCCTCCAGCA	180 55	TCGACCCCAS GOOTCOOTOC TTTTCCACAG AAGGTTAGAC CTTCAAAACA AAGGTTAGAAC
	CCTCATGGTC GTGCTGTCCT CTCTGGAAGG CACGCCTCG CGGAGCCTG TTGACAAGGA	340	ACCACCTATIG CARCITICATO PRINCIPLES ASSOCIATION ASSOCIATION
	CITICIACTICE GAGITITICAÇA ACAAGACCAC AGGGAAGATIG GAGAACTACS AGCTGATICCA	300	CECEREMENT CHARACTER CONTRACTOR CONTRACTOR CHARACTER CHARACTER CONTRACTOR CON
99	CICTAGROGO GICAAGITTA CCTACCCCAG TGAGGAGGAG ATTGGGGAACC TGACGTTCAC	09	CALABITATI GAZATICIOS GAGADICAGI CACTITICOCI GIAAATATOC

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8 55 TCCAAGAAAT CAAAAGAGCC ATCCAAGCTA AGGACACCTT CCCAAATGTC ACTATCCTGT ACAACCACGO TCTCAGGAGA TOTCTGATTT CCACAGACAT GCACCATATA GAAGAGAGTT TGAAGTTACA GTGTGTTTCC CTTTGGCTCC TGGGTACAAT ACTGATATTG TGCTCAGTAG AAATTCOOCA CAGCTGAGAG GAGACACAAG GAGCAGCCCG CAAGCACCAA GTGAGAGGCA 240 180 120 60

8

NTACTAMAA TACGAMATTA GCCAGGTGTG GTGGCACACA TCTGTAGTCC CAG

3173 3120 3060 3000 2940 2880 2820 2760 2700 2640 2580

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GAAGCCKGCT GGGGTGGTGG CTCACGCCTG TAATCCCAAC ACTTTGGGAG GCCAAGGCAG

OCOGATICACO TGAGOTICAGO ARTICOAGAT TARTICTOCCO AACATGOTIGA AACCOCCATIVI

CACIGCIGIT TITICCICIT IGGICCITCI ATCACIAAAA CICAICICAI TCAGCCITAC TACCICITCI GINAAGCIIT CCCIGGINIC AGGAAICAAA AITNAICAGG GAICITITICA

AGCATAACTA ATTATTTOTT TTCCTCACTA CATTOTACAT GTGGGAATTA CAGATAAACG

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CTOTOCHDAT GCACCATOCA TOCTCACAGT CCCTTGCCTA TOTOTGGCAG AGTOTCCCAG

AAGTUAGAGA AGCATUAAAA ATGAGCAGGG GCCTOGATCA GTOGGGTGTA TTCAGAGCAC

CCAGATOTOT GCCCCCACCC CATOTCCATT TACATOTCCT TCAATGCCCA CCTCAAAAGG

8

TIGCAGCTISCA AGTAATAGCA TOAACAGTCA GAAAAATACC TTATOAGGGG GCAGGGCTGA GICIGCICIT GIGIAGCICA GGAGACAATI CCAGCACAGA CACIACAGII AACGCIGAAC

AGCTGGGCCT TGAAGGATGG ATGAAATTTG GATAGAGAAT GAGGAAGACA GAGGGCCTCC

5

CAACTACCAT TAGCACTATG TTAGGAGCTG CAAGGCCCCA AAGTAGAAGA TGTGCATAAT

2460 2400 2340 2280

2520

TGCAATTAGG CAGATAAAGA CATCAGTCCC AGTAAATGAA TCCATAGACT CATCTAGCAC AAGTATAATC GGCTAACTCC TAAATCCCCAA TGAATAGTCC TAGGCTOGAC AGCAATGGGC TCACAACTGA GCAAGACATT CATATGATCA TITTAAGGAAG TOTTTCCCTT ATGTOTTJAGC ACAAACAGAT CACCAGCCAG CTTACACAGG CATTAACTCT CCTCAATGAG GAAGAATCAT

0

S

CAMAGACTCA GAGAACTAGA GITTAAGCIG AGGCAGAGIG CCGCCACCCT GGCATGCCCC

2220 2160 2100 2040

ICTCCAGAAG GAACITIOGGA GATGATIOGTO CAGAITGATIGA AACTIOOGTTC ATCCCAGITIC CTOGOGRACT GGGTATAATC CAACCATCAA AATAGAAGAC CTTOCAAGAA GCAGAGTCAT

3 2) INFORMATION FOR SEQ ID NO: 175:

50 Ê SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 175: (A) LENGTH: 991 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

8 55 50 45 6 CAGTCAGTAT GIGIGAAGAT CCCTGGTGCG TGGCCTTCAC CACGCATCTT GAGCAAATTA GCAGGGTCTA GAGACTOCTG GGACACTTTT CTTGGAGTGC TACTTCAGAA GCCTTATAGG TAATATTTT ATGCCACACT GGGATAAACA AGCAAGATTG CTCACTTCTG GAAGCTGCAT AGCCATGCCC AGTATTCCCA CTCTCCAAAA GGAACTGACC AGCTTATATT TCTCACACTT GCATAACAGG CTTAGTAAGT CCAAACACAG ATGACAGTGC TGTGTGGGTC TCTGTCAGAG AGCAGAATGT GGGCTGCATA TAAGCACACT CATCCCTTTG TCTGGGAATC TTTGTGCAGG GGAAAANGTA CCCTICGCTI GAGGCAGANG CAGCCCTICC CCCGAGNGCA NGGCINGGAG GGATCAATAT TITGCACACC TGTAATAGGC CATGOCACAC CAGCCAAGAT GCTCTGCTCA ATGACTAGAG AAAGCAGGTT ACCTAGTATA GTTTTTCCCAA ACTTCTTCCC ATCATAGCAC ATGTAGAAA titotogetet cagecatota gacacaetet ceaaatogag tottogaaaa tottettiet ATTITICITIC TOGCCAAGAT TICCITCIGT ATCACICCAA GCAGCCTCAG CAGAAGAAC GCCTCTTGTG ACTGGAGGTA ACAACCCTGC CCAGTAACTG TGGGAGAAGG 1980 1920 1860 1800 1740 1680 1620 1560 1500 1440 1380 1320

OCCTANATOG ACADATOGAT OCATACCCTT CCTGANATGA CTCCCTTCTG ANTGANTOAC CANACISCEA CATCICAGEC ISTANGENAA GEAGGAAACC TICTGETIGG CATAGETIGI CCANTACTOG CAGGITCCCT GGATCCAGAT CITICICITGCC CAACTCITAC TOOGAGATTG ATGAAATTGT GATCTAGGCT GCTGGGCTGA ATTCTCCCTC TGGAAACTGA GTTACAACCA TIOCIGATAA GATGOOGAAA GCCAGCACAC AGGACAGTAA ACCICCIGOO ACTICAAGCI AGTOCANGGT GCTTCCCTCC AAGGAAGAGC CAGTGAACAC AGTTTATTCC GAAGTGCAGT ATATCATOGC TICAAGGAAC ACCCAGCCAG CAGAGICCAG AAICTAIGAI GAAAICCIGC 1140 1260 1200 1080 1020 960 90

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20 TITITOTICCO TITITICAAG AGAAGACAAG AIGCIGCCIC AAAGAAAACC ATATACACAT 840

COGGSTIGGT GAGCGIGGTG GCTAIGTICT TICTGCTTGT TCTCATTGTG TCTTCAGTGT

ACTECATETE TOCCEOGRAG CICTOTORAG ACATEGRAAT GOGETTEEGT ACTEACEACA 780 720

CTGAGGACCA AGAGCTGACT TACACGTGTA CAGCCCAGAA CCCTGTCAGC AACAATTCTG

15

TGACATACAA TIGGAGICCC CIGGGAGAAG AGGGIAAIGI CCIICAAAIC TICCAGACIC 660 600

CTGTGAACAG CACCTGTAAT GTCACACTGA CATGCTCTGT AGAGAAAGAA GAAAAGAATG

540 480 420 360 300 240

ACAACCTGCA AATCTATCGT CGGCTTGGGA AACCAAAAAT TACACAGAGT TTAATGGCAT CAGGAGACTA CAAAGCAGAC ATAAATACAC AGGCTGATCC CTACACCACC ACCAAGCGCT

5

GGATACATIGC CITAGGICCG AACTACAATC TOGTCATTAG CGATCTGAGG ATGGAAGACG

CACCAGGAGA CTCAGAAACA GCACCCGTAG TTACTGTGAC CCACAGAAAT TATTATGAAC AAGAACCACG GCAAGTIAAA ATCATTOCTT GGACTTCTAA AACATCTGTT GCTTATOTAA

	WO 98/39448	PCT/US98/04493	WO	WO 98/39448	PCT/US98/04493	
	405			400	•	
			4	ACCTOGAGGC CCAGCAGCAC AACCACCATA GCCGGCCTCA GGGTCACAGA AAGCAAAGGG	099	
	COLCATIGGA GACTOTICCAG ATCATTAAGO COTTAGATOT GTGOTICGAGA ACCAAGAACO	300	O	CACTCAGAAT CATGGCACCT AAGTCTGGAC ACTGCCATCA GGGTTGCATT GGCTGTCGCT	720	
2		360	8	GTGCTCAAA CTGTCATTT GGGACTGCTG TGCCTCCTCC TCTGTGGTGG AGGAGAAGA	780	
	TGAGAAAAT CAGCAGCATT GCCAACTCTT TCCTCTACAT GCAGAAAACT CTGCGGCAAT	420	4	ANGSTAGCAG GOCGCCAAGC AGTGACTTCT GACCAACAGA GTGTGGGGAG AAGGGATGTG	840	
;		480		TATTAGCCC GGAGGACGTG ATGTGAGACC CGCTTGTGAG TCCTCCACAC TCGTTCCCA	006	•
0	ATGACAACTA TGATCAGCTG GAGGTCCACG CTGCTGCCAT TAAATCCCTG GGAGAGCTCG	540	2	ITGGCAAGAT ACATGGAGAG CACCCTGAGG ACCTTTAAAA GGCAAAGCCG CAAGGCAGAA	096	
	ACCIPITICE ACCEPTANT ANDARATIC AFGARCHASE GEOTICAGOS TGATGACAAG	009	8	GAGGCTGGG TCCCTGAATC ACCGACTGGA GGAGAGTTAC CTACAAGAGC CTTCATCCAG	1020	
15	GAACCTETAT AGTGATCCAG GGATGAACAC CCCCTGTGGG GTTTACTGTG GGAGACAGCC	099	15	GACCATCCAC ACTICCAATGA TATAGGAATG AGGTETGAAC TCCACTGAAT TANACCACTG	1080	
!	CACCTTGAAG GGGAAGGAGA TGGGGAAGGC CCCTTGCAGC TGAAAGTCCC ACTGGCTGGC	720		OCATITIOGOG GCTOTITYAIT ATAGGAGIOC AAAGAGTITCC TITTATCCICC CCAAGGAITG	1140	
	CTCAGGCTGT CTTATTCCGC TTGAAAATAG CCAAAAAGTC TACTGTGGTA TTTGTAATAA	780	ā	ABARTACAAT TRAITITITICT TACCARACA COCTITITION CITYCHOLD TRAITITION	000	
70	· ACTICIATUTG CITGAAAGGGC CITGCAGGCCA TUCTTGGGAGT AAAGGGCTTGC CITTUCCATUT	840	: ¥ 23	OCCUPATION OF THE PROPERTY OF	7,000	
	AATTTATTOT GAAGTCATAT AGTCCATGTC TOTGATGTGA GCCAAGTGAT ATCCTGTAGT	006	<b>:</b> }	ALTHURAN MANAGEMENT AND AND AND AND AND AND AND AND AND AND	0071	
75	ACACATTOTA CTGAOTOOTT TITICTGAATA AATTCCATAT TITIACCIAAA AAAAAAAAA	096	, ,	TOUR TRANSPORT VARIABLE TO THE TOUR TRANSPORT TO THE TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TOUR TRANSPORT TO THE TRANSPORT TO THE TOUR TRANSPORT TO THE TRANSPORT TO THE TRANSP	0621	
3	AAAAACTCGA GGGGGGCCC GTACCCAATT T	166	}			
			8	(2) INFORMATION FOR SEQ ID NO: 177:		
30	(2) INFORMATION FOR SEQ ID NO: 176:		30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2290 base pairs (B) TYPE: nucleic acid		
96			35			
3	(B) TYPE: nucleic acid (C) STRANDENESS: double	•	3	(xi) sequence description: seq id no: 177:		
	(D) TOPOLOGY: linear		ğ	TOGGCCCCT TITICARIECT CTOGGTOTT TICCCAAGAG TIACAGAIG TCAAGIGIGG	09	
4	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 176:	4	40	GENGETCHAC ACCEPTACTS TOGACCHATS ANGRESTITE CAGACCAGST GETTECHAAC	120	
	ACAGOCOTOT TOGGAGOCOTG AGOCOGGOTC TOCOTCACTOR COTCAACOCC CAGGOGGOO	09	A	ATTTCCAGGC TCCAGGAGAG AGCTGGGAG CCCCACAGA AAGCACAGGA AAATGCAAAA	180	
;	CTCCACAGGG CCCCTCTCCT GCCTGGACGG CTCTGCTGGT CTCCCCGTCC CCTGGAGAAG	120		AAAAAACAOT CTITTITITI TITITIGCITI TIAITATGAA AACAAACAA ATOCCCCAGG	240	
3	AACAAGGCCA TGGGTCGGCC CCTGCTGCTG CCCCTRCTGC YCCTGCTGCM GCCGCCAGCA	180 4	Ct SA	AGAAGGGTCC ATGATTACCA GAAACATCAA AGAGTACTTT CTACCATTTT TATTCTGTTG	300	
	TITCTGCAGC CTROTGGCTC CACAGGATCT GGTCCAAGCT ACCTITATGG GGTCACTCAA	240	2	TOTTGAGGCC AGCATTGCAA TAAACAAGCT AAACTACTTA CATTGGACTC ATTTTCAGTA	360	
20	CCANACACC TOTOLOCOTO CATGOSTOSC TOTOTOGANA TOCOCITICIO CITICIATIAC	300	50 AC	actgacattt acaggaatat actagaaacg gcactaaaaa gtttaagaaa agttacggta	420	
	CCCTGGGAGT TAGCLAYAGY TCCCRACGTG AGAATATCCT GGAGACGGGG CCACTTCCAC	360	\$	AACTIGGAIG CACAICAIAC AGAAAAGIAA CATTITAAAT ATAAAAAAGA AAAACTICCT	480	
;	GOCAGTECT TETACAGCAC AAGGCGGCT TECATTEACA AGGATTATGT GAACCGGCTE	450	. S	GGAAGCATTA TOCCAGTATT AAGGAACAGT GCTACTCTGG ATGTGACAAA TTCTGTATGT	540	
55	TITCTGAACT GGACAGAGGG TCAGGAGAGC GCCTTCCTCA GGATCTCAAA CCTGCGGAAG	480		GGGTGTTACT CTTTCCCAAA AGACTGTCAG AGGCGTGAGT GCTGCAAAAG AACAACAACA	. 009	
	GAGGACCAGT CTOTGTATTT CTGCCGAGTC GAGCTGGACA CCCGGAGATC AGGGAGGCAG	540	\$	nanachala cachanana tototettae actitotrag caagatgaca ctececaaca	999	
9	CAGTIGGAGT CCATCAAGGG GACCAAACTC ACCATCACCC AGGCTGTCAC AACCACCACC	. 009	<b>3</b>	CANAGAGGG TCTGGAGTTC AGTTCAGGCC GGAAGCCTGC CCCCTCGGCC TCCAGGGGTC	: 720	

8 5 8 33 30 25 20 2 5 S CCTCCACCCC TOGGGAGGGC AGACAGGCTC GGGARGGCCT GGCCAGGCCA CTGGAGGCTG ACAAAGTTTA TATTATATAA CTGGGGTTCC CTAAATTGAT TTCTTTTAAA ACAGTCTTAA GCAGOGAGCA GGCATGTCCA CCCGCAAGCC TGGGAGGCTA ACTCTGGCAT TCCTGGCCGG GCGTGAOCAG GGAGCACCGT GCGAGTCTCC GGGAGGGAAAT CCTCCTGGGG CCCAGAGACT CCCACGCTGA GGTCCGGGAG AATGCCTGGT TTCAGTCATT TCCGGACTAA CTGTGACAAC ACCIACAICI GIAAGOGICA GIOGACICIG AAICAAIIII AIOGIIGIII TAAAAICACC ACTTAATTAA AAGTTTAACA AACTGGCTGA AAACTCACCA AGTGTCAGAC TCACCAGCAA CGAGGGTAAC TITAAAAAAT GGAAACTITIC AAATCCATIT ATATTITIAT TATAAACAAA CIGIOCIOCO AGCACGITAC CAACCAGCCI GCGIGAAGAC CIGICAACIG ICGIGIGIGA GOCCOCTCCC CCGACGOCTC ACACAGOCAG CACCTCACTG CCCTGTGGCT GGAGGGGCAT CAGAGAACCA GCCGAGAAGG AAAGGCCCCA CGATGCTCCC TGTGCGCTGC CCCCACAGCC CAGAAAGCCT TAAAAAAAGTG ACAGCACCAA TGCAGCTGCT CAGTGTACCC NCCGTGGGCT TGAAAGCTGT AGCCTCTGGG AAAACAAAAC CAAAACATCA CCTTCTATTA AACTCTGTAT ATTATTATTT AGCCGCCATG CTCATTGGTG GGCCAGTTTG GGACATCCCC GTACTCAAAG ACCATATGGC GECCACCATA CAGGACAGAC CACACCACAG CTCCATACCC AGCGTCTGCC TGGAGGCTCC GTGTATTAGG ATACTAATGA TAGTCCCTAT ATCCATCCAG AAATGCTGGC AGAAAGCACT TTTAAAAAAT GATAATTTAC CAGCATCICC TCATCAGAGT TCCCTCTCCA GTAAGGGTAT ATTOCTTAAA TIOGGITIAA ATAGIOCATT AAAGATOTGT TTAGAAAATA COTTTGAAAA TOCAMOUNDO OCCCCCAMO CCCAMOGCACO CCCGGCTTAG GOTOTACIGTA TCACCCAGCO GTCAGGGTCA GTGGCTTCTT TCTAGATGAA AGGAGCAGAG GCGAGCCGAC GCCACCGTCA GCGCTGAGGA AGTCATTAAT CCTTCGAAAC TCTGAAAAGA AACCAGTGTT GAAGTCTGGA CCGCAGCCTT CTAATACAGA AGAAACGGAC GTGACTGTCA CCCTCAGCCC GCCAGCAAGG ITTACAATAG AAAGITAAAA AICAAGACIT AGAITTACTA TACAITTITIT CICICAGATT NGAGACCAGA AGTGAATACA AAAGAACTAA ACAAAATAAA AAATTAGAAT GTGCTGTAGC NAAAAAGCAT GTTAGAAGCT GCCCTACAGG TCTCAGCAGT GGGACAATCT AATTGAATCA ATTICAGAGTG TICTCAAATC CAATTCCGAC ACACGACTTG TCACTACTCC TCTCCCCTTG 2280 2100 2040 1980 1920 1860 1800 1740 1680 1620 2290 2220 1560 1500 1440 1380 1320 1260 1200 1080 1020 960 900 840 780

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CCCAGAATTT GCAGTAGCTC AAATTAAGTT TCTAGCTATT AAAAAGAAAA

549 540 480 420 360 300 240 180 120

GAATGCTCCT TCACCAGGAC CAGAGAACTG ATTTACAGAA GTGACATGAA

aaaaaaaaaa AACATTCCAT 20

IGITICCAAG AIGCITCIGA AGATIGCCIA AAAATAGCCG GITICCACCC CCGIQAAIGC AGTAGAGAGG TAAAAGGCCA CCATCTCCTT GACCTCTGGG GAACTCATCC ACAAAGAAGA TIBSTECAGS AATSTAGTAG GCATACACST GSTTBCGTBG ATCTBGGCCC TCCTBATGTB ATCAGTAACA ACCAGGAGAG AAGCTGCTGG AACTGACCTC TGGGAACTCC CTGGGATGGT TOTOGRATAG TOTOTOCA TOCCTOTOCT CATOGGCTAC CACCTOTOCC ACCOTOGTTA CICAGCAGCI ACCIGCATIG IGGCCAAAGG AIGACCIAIT CCIICICAGG AGGGCAAAAA GECACGAGCC ATGCCTGGCC TCTCCTTGAT TCTTACAGTC ACTITIOTTGG CTGTTTCTGA

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3 INFORMATION FOR SEQ ID NO: 179:

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 1509 base pairs

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 179:

GCCCGCCGCA TCCGCCGCCGC CAGCCCCCAG CATGTCGGGC CCAGACGTCG AGACGCCGTC GGCACGAGGG CTCATTCATT CCGCGCCGGG CCTGCCAGAC ACCTGCGCCC TTCTGCAGCC 120 60

25

CAACAGCCAG AACGGGGAGC GCTGTGTGGC CGCCCTGGCT CGTGTCGAGC CGCCATCCAG ATCTGCCGGA TCATGCGGCC AGATGATGCC AACGTGGCCG GCAATGTCCJ COGGGGGACC ATCCTGAAGA TGATCGAGGA GGCAGGCGCC ATCATCAGCA CCCGGCATTO GCACCGACTI 180 300 240

GIOGRAGGICC AGGICARCGI GAIGICCGAA AACAICCICA CAGGIGCCAA 420

CCTGTCTCCC ATGTGCATCG GTGAGGTGGC GCATGTCAGC GCGGAGATCA CCTACACCTC

360

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CAAGCACTCT

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INFORMATION FOR (SEQ ID NO: 178:

AMBOURANCE AND MOROCCA COUNTRIESTA TOTOCCCCTG TOGCTGAMGA ATOTOGACAI 480

540

GCGGTATGAA GCCCAGAAGC TOGAGCOCAT GGAGACCAAG TOGAGGAACG GOGACATCOT GGTCCTCGAG GTGCCTCCTG TTGTGTATTC CCGGCANGAG CAGGAGGAGG AGGGCCGGAA

600

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WO 98/39448

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(A) LENGTH: 549 base pairs
(B) TYPE: nucleic acid
(C) STRANDENKESS: double
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO:

178:

•	WO 98/39448	PCT/US98/04493	W		PCT/US98/04493	
	409			410	<i>i</i>	
				TITICAGTOC COCACAATGA GTCAGAAGAT GAAGTOGCTG TTGACATGGA ATTTGCTAAG	480	
	CCAGCCAGTC CTCAACCCAG AGCCGAACAC TOTCAGCTAC AGCCAGTCCA GCTTGATCCA	099		AATATOTATG AACTGCATAA AAAGTTTCT, CCAAATGAGC TCATCCTGGG CIGGTACGCT	540	
S	CCTGGTGGGG CCTTCAGACT GCACCCTGCA CGCCTTTGTG CACGGAGGTG TGACCATGAA	720	S	ACCIONOCATO ACATCACACA GCACTCTOTO CTGINITCCAT CAGIACTACA GCCGAGAGGC	009	
	GCTCATGGAT GAGSTOSCOG GGATOSTOSC TGCACGCOCAC TGCAAGACCA ACATGGTCAC	780		CCCAACCC ATCCACCTCA CTGTGGACAC AAGTCTCCAG AAGGGCGGCA TGAGCATCAA		
	AGCTTCCGTG GACGCCATTA ATTTTCATGA CAAGATCAGA AAAGGCTGCG TCATCACCAT	. 840	:	AGCCTACOTC AGCACTITIAA TGGGAGTCCC TGGGAGGACC ATGGGAGTGA TOTTCACGCC	720	•
9	CTOGGGACOC ATGACCTTCA CGAGCAATAA GTCCATGGAG ATCGAGGTCT TGGTGGACGC	006	2	TUTGACAGTG AAATACGCGT ACTACGACAC TGAAGGCATC GGAGTTGACC TGATCATGAA	780	
	CONCETIVIT STOCKLAGET CTCAGANGOG CTACCOGGCC GCCAGTGCCT TETTCACETA	096		GACCIGCITY ACCCCAACA GAGIGATIGG ACTCICAAGT GACTTGCAGC AAGTAGGAGG	840	
15	COTISTICACTIO AGCCAGGIAAG GCAGGTCGCT GCCTGTGCCC CAGCTGGTGC CCGAGACCCA.	1020	15	GOCATCAGCT COCATCCAGG ATGCCCTCAG TACAGTOTTG CAATATGCAG AGGATGTACT	006	
	GGACGAGAAG AAGGCTTTG AGGAAGGCAA AGGGCGGTAC CTOCAGATGA AGGCGAAGCR	1080		OTCHORANAG STOTCAGCTG ACANTACTOT GOGCCOCTTC CTGATGAGCC TGGTTAACCA	096	
;		1140	Ş	AGTACCGAAA ATAGTICCCG ATGACTITIGA GACCATGCTC AACAGCAACA TCAATGACCT	1020	
8	GCCATGGCAA COGGCCAGT OTCCAGTCAC TTAGAAGTTC CCCCCTTGGC CAAAAACCCA	1200	3	TITGATGOTG ACCTACCTGG CCAACCTCAC ACAGTCACAG ATTGCACTCA ATGAAAACT	1080	
	ATTCACATTO AGACCTOOTG TIGICTGAAG TITTCOTATC ACAGTOTTAA CCTGTACTCT	1260	-	ISTANACCTS TGANTGGACC CCANGCAGTA CACTTGCTGG TCTAGGTATT AACCCCAGGA	1140	
25	CTCCTGGAAA CCTACACACC AAAGCTITAT TTATATCATT CCAGTATCAA TGCTACACAG	1320	25	CTCAGAAGTG AAGGAGAAT GGGTTTTTTG TGGTCTTGAG TCACACTGAG ATAGTCAGTT	1200	
	TOTTOTICCCS AGGECCOGGA GGCSTTGGGC AGAAACCCTC GGGAATGCTT CCGAGGACGC	1380	J	CIGTOTGACT CTAATAAACG GAGCCTACCT TTTGTAAATT AAAAAAAAA AAAAAACQ	1260	
	TOTAGGGTAT GGGAAGAACC CAGCACCACT AATAAAGCTG CTGCTTGGCT GGAAAAAAA	1440	-	SERGEGEGES CCCGOTCCCA TISSCCCTIT NOTABATICET NITACAATICE CTAITSC	911	
30	ADADADADA ABADADADA RADADADA BADADADA ABADADADA ABADADADA	1500	90			
	AGAAAAN	1509				
36			35	(2) INFORMATION FOR SEQ ID NO: 181:		
3	(2) INPORMATION FOR SED ID NO: 180:					
				(B) TYPE: nucleic acid		
9			6	TOPOLOGY: 1in		
	(B) TYPE: nucleic acid (C) STRANDENESS: double			(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 181:		
!	(D) TOPOLOGY: Linear		, G	GGCATGWKCA GACATGACTT CTATTGCCAG GCTGGTCAAG TGGCAGGGTC ATGAGGGAGA	09	
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 180:			CANCGATAAG GOTGCTCCTT ATGCTCCCTG CTCTGGAATC CACCAGCGGG CTATCTGGGT	120	
	ACCTOTATCA TAGGAAAGAT GOCCACACOS GOGGTACCAG TAAGTGCTCC TCCGGCCACG	09	F	TIATOGOGCT GOGGACTAGA ATTGGATGCT TCAAAACCAT CACCTGTTGG CCAACAAGTT	180	
20	CCAACCCCAG TOCCOGCOGC GOCCCAACC TCAGTTCCAG COCCAACGC AGCACGGCT	120	50 1	TGACCCAAAG GTAGANGATA ATGCTCTTCA GTGCTTAGAA GAATACCTAC GTTATAAGGG	240	
	GOGGCTCCGG TTCCCGCTGC GGCTCCAGCC TGCATCCTCA GACCCTGCGG CAGCAGGGGC	180	О	CCATTCTATT GCGACCTGAA CTTTGAAGAC CACAMTATTG AAGAGGCGTT GCTTACCYGT	300	
	TIGNACTIOCO OCTICCTIOSCE AGACCICOGO CTCAGCOCIAA NTOCIAGOGCA GACCOCIAGOS	240	-	TOOGOOCCAA GAGGCATGTT ACCAAACATG GYYCARGAAM YTTGGYKGGG AMCARYGGGKG	360	
55	CCCGCTCTGC CTGGTCCTGC TCTTCCAGGG CCCTTCCCCG GCGGCCGGT GGTCAGGCTG	300	С	GKKGGGARRM CHRGGGYTTG SCAAWTTCSK KGGCMCCYT TTAGGGTAAR RRGGGCKGTW	420	
	CACCCAGTCA TITTIGGOCTC CATTGTGGGC AGCTAGGAGA GAGGCAAGGA GGGTGCTGCC	360	×	ATTAGATTOT GOSTAAASTA GGATCTITITG CCCTTGCAAA TTTGCTGCCT GGGTGAATGY	480	
. 09	CGAOTTATOG GGACCCTOTT GGGAACTOTC GACAAACACT CAGTGGAGGT CACCAATTGC	420	6 6	TGCTTGTTCC TTCTCHACCC CTAACCCTAG TAGTTCCTCC ACTAACTTTC TCACTAAGTG	540	

WO 98/39448

PCT/US98/04493

Ξ SEQUENCE CHARACTERISTICS:

55 (2) INFORMATION FOR SEQ ID NO: 184:

50 AAAAAAAACT C 791 780

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TAAAAAAAA AAAAAAAAAA CTCGA

GATTOGTOTA AATCITTITTA TAAATACATA AATAAAAGNA AAATATGCAT TITITCTTTTC

TCATATATT AACTITGCAA AAAGATTTAC TTTGTACATG TTACAGGCTI

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1405

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TTTTGGATTT

GGGAAACTAA TIGIATITAG IGACAAAAAT AAAAAGIITI TITITIATAA

TICAGICIOC

1260

1200 1140 1080

CTOTACAGAC CATCTOTATG TTAGGTGACA TTGATTATGG GTTATAATCA TITATCTGAG TITAGTOGTC CTAATATATA TOTAGAGAAA GATOGTOGGG ATTTCCTTTA AATGGTAAGA GTTTCTAAAA CAGACAATAA TTTAACAAGC

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TITICIAGAC TIOGGATCIG CAAGAAGGCC AATTGCCTAA AATTICIGAG AACAGIGCAC CACCACGACC AACCATATGG CAAATGAACC AAGCCCAGTT GTTGCAGTGA TTGGTTGTCT

AAGATTATTT TATCACTACA AGCTTTTAAC TTTTTAAGTT ATTGTACAAG TATTCTACCT

1020 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 6

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TATATOSCTT TECTTCACCT CIGOGTCATG ATTOTTCTGT TGACTTACAC ACCAGAAATG

AGTATICGCC TOOGAATITIT ICICCGAAGA TACCCCATAG COCGAGITIT TOTAATTATA ACTAATCTGG CAGGAATGTA CGGAAAAGTT CGCAAAGCTG CTAGTTCAAT TGATCAGTTT ATTGACAATG GTGAAGGCAC TOOTCTGCGA AATGTTCCTG TTCTTTTTAA TGACACAGAA CAGCAGATGA ACTOCOCCTO TOGAAGTAGT AGTAATOGGT CTTOGATTAA TATOTOTOGA ATGCTGGAGA GTCTCAGCAC AGAAAAGAAC TCCCTGGTCT TTCAACTGGA GCGCCTCGAA CAGTOTOAGT TAGAAAATOG ACTOCATOAG CTAACAGAGA CTOTOATOCA GAAACAGACO GACGAAGAAA TTCAAAAACT CAGGAATCAG CTTACCAATA AAACTTTAAG CAATAGCAGT TATATAGAAG AAGATCTTTA TCGAACAAAG AACACATTGC AAAGCAGAAT TAAAGATCGA CAGAAAGCAT CCAAACAAGA ACTAGAGACA GAACTOGAGC GACTGAAGCA GGAGTTCCAC GTTAATGAAG CAGAATCAGC AAGAGAACAG TTACAGGWTC TGCATGACCA AATAGCTGGG AAGCTGATGG GCCAGATACA TCAGCTCAGA TCCGAATTAC AGGATATGGA GGCACAGCAA

45 8 3ဗ TOGITICITI CCICCATACC CCITCIGCAT TICAGIGITI TIGITIAGIT TICCIGGITT CCACTAATAG AATTCAGCTT TTAGCATGOG CTGTTTCATA CTGTTCTGAT GAAACTGATT CACCACACCT ACAGGGACCA COTOGTOGGC TOTOGACTAG COGCCAAGCT CCCTGCAGGC CCTGCACCAC CITGAAAGAC ATTICTAATA TGGTTIGICA GGCAAAGIGG TIAGIAGICA TOCCCCTOGA CTOGGCTCTG TOAGAGTOGC CTTCTGCACT GTGCACAGTA GGTGTGAACA TTTGTGGCCT GAGGTAGAAG TCCTCAGAAA TCAGCAGACT TCACTGATAA AATGCTGACT GATCYTGGGA AAGANATATC TIGCCAGGAA AAATGATAGN CCTIGACAAT GIIGAAITGAI ACATGTOGGC TOGAGATCTA TICATITICOT TITIGGCTIGA ATTITICIGRA TOGITTACIT TECGTGGECA GAACEECTEE AGGTEAGAGG CAGAAGAGA GEETEATIGG TEACAGEAGE GOTETITIAG ACTOTICITI TITICCCANCI TETEACETEC TOCCCCIECT TEAGGOTACT 720 660 600 540 480 420 360 300 240 180

25 ACTOGICTIC AAAIGIGIAC AIGIGIGCCA GGGAGCAAAI GCCIICTIGI TICIGAAAIT

GGCACAGATA ACTATOTACA TOTATTOCTT AAATGITTIT TIAAGITITA TAITCITGGC 120

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SEQUENCE DESCRIPTION: SEQ ID NO: 182: (A) LENGTH: 791 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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3 INFORMATION FOR SEQ ID NO: 182: Ξ SEQUENCE CHARACTERISTICS:

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777 720

AGGITITGGIA GCGITGGAGGA GAACITITGAI GGAAAGAGAA CCITICCCIIC IGITACIGIIA TCAGTIGAAT GCCIGCIGGT AGCITTICCA TICIGIGGAG CIGCCGIICC TAADAATICC

660 600

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AAATTGATTA ACAGCTTGAA AGAAGGCTCT GGTTTTGAAG GCCTAGATAG CAGCACTGCC agtagcatog agctogaaga acttcggcat gagaaagaga tgcagaggga ggaaatacag

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 183

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agaatgagaa ctoctotoat agogagagto aaggagggat atotogtaga gcacttgatt

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3

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1405 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 183:

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414

. 8 120 180 240 300 360 420 480 540 9 999 720 780

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linear	
LOGY	
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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 184;

~	STCATISCAST COCCOGRAGA ACTOTOCTOT TTGAGGCCGA COCTAGGGGC CCGGAAGGGA	09
	AACTOCGAOG CGAAGGTGAC CGGGGACCGA GCATTTCAGA TCTGCTCGGT AGACCTGGTG	120
01	CACCACCACC ATGITGGCTG CAAGGCTGGT GTGTCTCCGG ACACTACCTT CTAGGGTTTT	180
2	CCACCCAGCT TTCACCAAGG CCTCCCCTGT TGTGAAGAAT TCCATCACGA AGAATCAATG	240
	GCTGTTAACA CCTNGCAGGG AATATGCCAC CAAAACAAGA ATTGGGATCC GGCGTGGGAG	300
15	AACTGGCCAA GAACTCAAAG AGGCAGCATT GGAACCATCG ATGGAAAAAA TATTTAAAAT	360
	TGATCAGATG GGAAGATGGT TTGTTGCTGG AGGGCTGCT GTTGGTCTTG GAGCATTGTG	420
Š	CTACTATIGGC TIGGGACTOT CTAATGAGAT TGGAGCTATT GAAAAGGCTG TAATTTGGCC	480
3	TCAGTATGTC AAGGATAGAA TTCATTCCAC CTATATGTAC TTAGCAGGGA GTATTGGTTT	540
	AACAGCTTTG TCTGCCATAG CAATCAGCAG AACGCCTGTT CTCATGAACT TCATGATGAG	009
25	AGCCTCTTGG GTGACAATTG GTGTGACCTT TGCAGCCATG GTTGGAGCTG GAATGCTGGT	099
	ACGATCAATA CCATATGACC AGAGCCCAGG CCCAAAGCAT CTTGCTTGGT TGCTACATTC	720
Ş	TGGTGTGATG GGTGCAGTGG TGGCTCCTCT GACAATATTA GGGGGTCCTC TTCTCARCAG	780
3	AGCTGCATGG TACACAGCTG GCATTGTGGG AGGCCTCTCC ACTGTGGCCA TGTGTGGGCC	840
	CAGTGANANG TITICTGANCA 1933/1920.CC CC1939A/FIG G3CC1933/TC 1051/C11107	006
35	GICCICATIO GGATCIAIGI TICITICCACC TACCACCGIG GCIGGIGCCA CICITIACIC	096
	AGTOGCAATO TACGGTGGAT TAGTTCTTTT CAGCATGTTC CTTCTGTATG ATACCCAGAA	1020
. 6	AGTAATCAAG CGTGCAGAAG TATCACCAAT GTATGGAGTT CAAAAATATG ATCCCATTAA	1080
?	CTCGATOCTG AGTATCTACA TGGATACATT AAATATATTT ATGGGAGTTG CAACTATGCT	1140
	GOCAACTIGGA GOCAACAGAA AGAAATGAAG TGACTICAGCT TCTGOCTTCT CTGCTACATC	1200
45	AAATATCTTG TTTAATGGGG CAGATATGCA TTAAATAGTT TGTACAAGCA GCTTTCGTTG	1260
	ANOTITINGIA GATAAGAANC ATOTCATCAT ATTINAANOT TCCGGTAANG TGANGCCTCA	1320
ç	GGICTGCCTT TITTTCTGGA GAATAANTGC AGTAATCCTC TCCCAAATAA GCACACACAT	1380
3	TITICAATICT CATGITIGAG TGATITIAAA ATGITITIGGT GAAIGIGAAA ACTAAAGITI	1440
	GIGICAIGAG AAIGIAAGIC TITITICIAC TITIAAAITIT AGHAGGITCA CIGAGHAACT	1500
22	AAAAITTAGC AAACCIGIGI TIGCATAITT TITIKGGAGIG CAGBITAWIG TAAITARAGC	1560
	ATTCCAGTAA NAGTGTNITT AAAGTTGNIC TATAIN	1596

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GOGCAGAGCC CGYACGAGCA GGACGACGAC GACAAGGGG ACTOCAAGGA AACGOGGCTG ACCCTGATGG AGGAAGTGCT CCTGCTGGGC CTCAAGGACC GCGARGGTTA CACATCATTT TGGAATGACT GTATATCATC TGGATTACGT GGCTGTATGT TAATTGAATT AGCATTGAGA GGAAGGTTAC AACTAGAGGC TTGTGGAATG AGACGTAAAA GTCTATTAAC AAGAAAGGTA TOGAATCCAT TAAAATTGCA TTATCAGTTA AGAAATGTAC GGGAACGATT AGCTAAAAAC ATCTGTAAGT CAGATGCTCC AACAGGGGAT GTTCTTCTTG ATGAAGCTCT GAAGCATGTT CITICITICACA AATGGGTGAA TGACCCTCAC CGCATGGACA GGGGCTTGCT GGCCCTCATT AAGGAAACTC AGCCTCCAGA AACGGTCCAG AACTGGATTG AATTACTTAG TGGTGAGACA ACACATCCCC TCACCAATAA CAACATTAAG CAGCGCTTCA TCAAGAAAGT ACAGGAAGCC CTGAAGGCCA ACACCAATGA GGTTCTGTGG GCGGTGGG CGGCGTTCAC CAAGTAACTC TICTIGGITIT CTGTAATITIG TACTITICCCA CACTATAATT GGCTTCTGTT TTACAAAATG TACCTIGACTE ATGCCTEGGA CGTECTIGGAG AATGCTTTTG CTCCTCTTCT GGACGAGCAG TAIGATITIGG CTACCAAGAG AGTGCGGCAG CTTCTCCAAT TAGACCCTGA AGTGGAATGT IGCTCGGGGT GAACCATTCT CCTTTCTCTC AAGTAAACCA GTAGTTTTTC TTCTGTTGAC CITICIGITY TYAAAAAAG TYTYACIGCC ATATYGGCAT TCCATYCCCT GIYGCCATCC STGGOTGCT ITTICITITY IGFACGISTA CAGGATICIG CIGGFACGAG AGGCCTTCCT TCACTOTTAC CTGTTTTGGG TTTCTGGTCT ACTTTGACTT TCAAAGTACC TCCAGCCTCC TCATAGGCAC AGCTTTTGGA TGACCTCAGC TTGAGTTTCT CCATATGTGC ATGTACATCT AGCATTCTGC CTACAGTTCA GACAGAAGTC ACAAAAAGGC CTTCAACTCA CCAAAGGTAA ATATICTOTAT CTATTAGGAC ATTITITACA TAGACTICAG ITGAGATGTA TACTTAGGA AATTAITITT AAATTGAAAC AGCACAGTAA ATACTTAATA TAAAATGTCC CTTGGATTTT GCTTCCCATG TAAATCTATT GTATTATAC ACTTGTTATA ATTTTAACTA TAAAGGTCCA ATTOTTICAC AGAGCCAGIT TGGGATGGGC TGCATTCCAT TTATGCTGTA TATAGITTGA SEQUENCE DESCRIPTION: SEQ ID NO: 185: (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2293 base pairs
(B) TYPE: nucleic acid
(C) STRANDERNESS: double
(D) TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 185: (X 2 5 2 9 25 35 <del>\$</del> 45 S 55

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960 1020 1140

1200 1260

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ATTATATATA AATTACCCCT TCTTCTGGCC ACCCTGCTC CCATCTTAGT ATTTTGCAAG

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330 25 20 15 5 S 2 AAAAAAAAAA AAA AAAAGTACTA GATGGTGTAT AACTCTAGAG TIGAATTITTA AGGGATTCCC TAATATGTAT GTTAATAGTG GTTTTAATTT TTTATATACA ATTAGGATAA TTAGCATTGT CAGACTATAA ACCTITGCTT TICCIRCCAA CIRATGATIA CAACCCAGAG GATIAAGAAT TITCIAACAG AAACCICTRI TACTCACACA TICTIGGGGA AAAAAAICAA AIGICAGICC TAGCAGAIGT TGCAIGTAAA TCTTGAGTTT TAAATTITAT CITACIGITT TIATAATTIC TATIOGGAAG AGGCTIGIGA CCAGTACCAA THITTGAGCT GIGCCAPTIA TOSTACICIT TOCCTATGCA TCCCCTITIT AGATTITITIT ACTATOTITT TATOTGAAGT AATAAATAAA CAATGATOTT GAAAGTOOOY RAAAMAAAA TTTAAAGTTT AAATCGACCT CTTGTCGCTG AAGGAGAGAG AAAAGATGTG TGTCTGATTG GTCCTGGGAT ATCTAATCAG TTGTACACCT GGTGCCCCTC GCTTGCTTCA ATCATGGTTA TTTGATGGCA INFORMATION FOR SEQ ID NO: 186: GGGCTTTCAG CTATATOTTO AACAAATTAA ATGTCAAAAT TITTTATTAC CATAGTCCAT ATTITTACTA CHIMINGIGE GIGITIAGAG ATTITITITIO TITICITIATIC ACTITATION ATCARCACCA TROOFFICAT CCACAAGTAA ATTAATATCT GCTCTGAAAT GTCATTTATC TIGITOTIAA CATICATIOC

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 1212 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 186:

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CATTITIGGCT GAAGGICCTA ACCIGATTAT AAAACAACCA GATGAGITGC TGGACAGCAI

420 360

STCAGATTOG TOTAAGGAGC ATCACOGGAA GTCTGGCCTT ACCAAGGCAG TGAAGGAGAG

TACAATTACA TIGCAGCAGG CAGAGIAIGA ATTICIOICC TITOTACGAC AGCAGACICC

540

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ATCCTACTCC ATGGGATGAA GACGTCTGAC ACTATTATCC GGGAGGGCAC CCTGATGGGC

720 660 600

CCCCAGAAAA TIGCAGGIGA ACICIAIGGA CCICICAIGC IGGICIICAC ICIGGIIGCI

CCTGCTCAGG TOCGAACAGG GCTCCTOGAG TCCATGATCC CTATCAAGAT GGTCAACTTC

480

GECCCCAGGG GAGAGCATGG CTCAGCCGAT GGTCTGGGTG GACCTGGAGA TGACAGGATT

GGACATIGAG AAGGACCAGA TIATIGAGAT GGCCIOTCIG ATAACIGACT CIGATCICAA

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CHCHACCICG

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CCGTGGGTTT GCGACGTTTA GCGACTATTG CGCCTGCGCC ACGCCGGCTG CGAGACTGGG

120

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GOCACGAGGC GAGCCGGCGC ACCGTACGCT GGGACGTGTG GTTTCAGCTC GTGCGCCTCC

GOOGRACIO CIOGROCCO GRANISCIAG GOGGCICCCI GOGCICCAGO CIGITIGOGG

GAGTCACGGA CGGTTCGGGG CCCGAGGTGT CCGCGAAGGT GGCGCACATG

240 180

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8 S 8 5 CCCACCICIC CICGACCCIG GACGICIACC INCCGGAGGC CCACATCITG CCCACICCGC GCTTCCGGAA GTTGCTTTTG TCCAAACATC CGGGCTTCTC CTTTTTGTGT GCAGCTGATG CAGCTGCTGC TGAAGAGGAG GATGGAGAGT TCCTGGGCAT GAAGGGCTTT AGCTICGAGG ATATIGGGTGA GCTGCATICAG CGCCTGCGCG AGGAAGAAGT ATCCAGGGG GAGGCTCAGC TGTGATTGAC ATGGAGAACA TGGATGATAC CTCAGGCTCT GOSCOGCGG GCGGCGCCG AMATGGAGCT GGCCCGGMAT GGGGAGGGTT CGAMGAAMAC GCGCGGGGCT AGCGCGGGTT TCAGCGACGG GAGCCCTCAA GGGACATGGC AACTACAGCG TOCAGGGCOT TOAGCITISTA CGCCAACATC GACATOCTICA GACOCTACIT TGATGTGGAG AMOGGACAGO TGAGOCGGCA GOTGGCAGAT CAGATOTOGO AGOCTGGGAA AAGACAAGOO AGACGCTGAT TCCGGCCGAT 240 540 480 420 360 300 180 120 6

2 INFORMATION FOR SEQ ID NO: 187: 2280

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2160 2100 2040 1980 1920

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1740 1680 1620 1560

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30 Ξ SEQUENCE CHARACTERISTICS: (A) LENOTH: 1605 base pairs

3E SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid (C) STRANDEINESS: double (D) TOPOLOGY: linear 187:

CTCCAGATTG ATTACTCAAG CAGACAGCAC ACGAAATACT ATTTTTCTCC TAATATGCTG aaaaaaaaaa an TATATISCATT TOCTTTTAAA CCATTTCTTT TOTTTAAATA AATAAATAAG TAAATAAAGC TITICCATTAT GACACAGCAG CICCITIGIA AGTACCAGGI CATGICCATC CCITIGGIACA TOTOGIOGIT TITITITICIC ACCORGANGS CITOGCAGAG CACCITOGGI TAACITGCAT GAMGACCOTO ACTIGATOCC ACTIATOATO CIOCCACIAC ATOCITATOI GGAGGOAACT CATCTTCAAG AAAAAAATAG ATGAAAAGAA GAGGAAAATT ATAGAAAATG GGGAAAATGA TCATAGGGCA CTTGATGACA TTAGTGAAAG CATCAAAGAG CTTCAGTTTT ACCGAAATAA AGAACTOTGC AGACGCTGOT ATCCAGAAGA ATATGAATTT GCACCAAAGA ATACATICCC CAGITICATGA AACATCITCA TTATAGAATA ATIGATIGIGA GCACIGITAA ICCAGGGCIC TGTCCACTTG CAGGAAATTC AGTTCATGAA GATAAGAAGT TTCTTGACAA AGGCTGCTTC 1212 1200 1140 1080 1020 960 900 840 660 780 720 8

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	ACAGCCATTG GCACCTGCTT CGGCTACTGG CTGGGAGTCT CATCCTTCAT TTACTTCCTT	780
δ.	OCCIACCIOT GCAAGGCCCA GATCACCAIG CIGCAGAIGT IGGCACIGCT GGGCTAIGGC	840
	CTCTITIOGGC ATTGCATTGT CCTGTTCATC ACCTATIANTA TCCACCTCCA CGCCCTCTTC	006
	TACCICITICT GACIGITIGGT GGGTGGACTG TCCACACTGC GCATGGTAGC AGTGTTGGTG	. 096
01	TCTCCGAACCG TGGGCCCCAAC ACAGCGGCTG CTCCTCTGTG GCACCCTGGC TGCCCTAAAC	1020
	ATGCTCTTCC TGCTCTATCT GCATTTTGCC TACCACAAAG TGGTAGAGGG GATCCTGGAC	1080
7	АСАСТОСАЛОС ССССЕЛАСАТ ССОЗССЕЛТС САВЛЕВЕТС ССАВЛЕЛСЯТ СССТОССАТО	1140
2	CTCCTGCTG CTCGGCTTCC CACCACGGTC CTCAACGCCA CAGCCAAACC TGTTGGGGTG	1200
	ACCTIGART CACACTGACC CCACCTGAAA TTCTTGGCCA GTCCTCTTTC CCGCAGCTGC	1260
70	AGAGAGGAGG AAGACTATTA AAGGACAGTC CTGATGACAT GTTTCGTAGA TGGGGTTTGC	1320
	AGCTGCCACT GAGCTGTAGC TGCGTAAGTA CCTCCTTGAT GCNTGTCGGC ACTTCTGAAA	1380
25	GOCACAAGGC CAAGAACTCC TGGCGAGGAC TGCAAGGCTC TGGAGGCAAT GCAGAAAATG	1440
}	GOTCAGCTCC TITGAGAACC CCTCCCCACC TACCCCTTCC TTCCTTTTA TCTCTCCAC	1500
	attotettec taartataga cittgotaatt aaaatottea ttegageete gaaaaaaaa	1560
30	aaaaaaaa aaaaaaaa aaaaaaaaa aaaaaaaac togag	1605
35	(2) INFORMATION FOR SEQ ID NO: 188:	
40	(1) SEQUENCE CHARACTERISCS:  (A) LENGTH: 1516 base pairs  (B) TYPE: nucleic acid  (C) STRANDERNESS: double  (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 188:	
45	ATTOGGCATG AGGGGGTCAC GTGGTGGCTG GGCGGGGAA ATGGCGGCTT CAGGAGAGAG	09
	COGGACTICA GOCGOCGAG GCAGCACCA GCAAGCAITI AIGACCITCI ACAGIGAGGI	120
ç	GAAACAAATA GAGAAGAGA ACTCGGTTCT AACTTCGAAA AATCAGATTG AAAGACTGAC	180
<b>)</b>	CCGTCCTGGT TCCTCTTACT TCAATTTGAA CCCATTTGAG GTTCTTCAGA TAGATCCTGA	240
	AGTTACAGAT GAAGAAATAA AAAAGAGGTT TCGGCAGTTA TCCATCTTGG TGCATCCTGA	300
55	caadaatcaa gatgatgctg acagagcaca aaaggctttt gaagctgtgg acaaagctta	360
	CAAGTIGCTA CTOGATCAGG AGCAAAAGAA GAGGGCCCTG GATGTAATTC AGGCAQGAAA	420
9	AGAATACGTG GAACACACTG TGAAAGAGGG AAAAAACAA TTAAAGAAGG AAGGAAAACC	480

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IACAATTIGTA GAGGAGGATG ATCCTGAGCT GTTCAAACAA GCTGTATATA AACAGACAAT GAAACTCTTT GCAGAGCTGG AAATTAAAAG GAAAGAGAGA GAAGCCAAAG AGATGCATGA AAGGAAACGA CAAAGGGAAG AAGAGATTGA AGCTCAAGAA AAAGCCAAAAC GGGAAAGAGA STGGCAGAAA AACTITIGAGG AAAGTCGAGA TGGTCGTGTG GACAGCTGGC GAAACTTCCA AGCCAATACG AAGGGGAAGA AAGAGAAGAA AAATCGGACC TTCCTGAGAC CACCGAAAGT AAAAATGGAG CAACGTGAGT GACCGCCAA GGTCACAGGC ACAGAACCTT TCCCCTGCTA TCTCCCTTCC TGCTTCGAAG GACTCATTCT TTCCTCCCAC TTCCACCCCA ACATAGAGTA GIATITIGCIT ITIAGICCAT ITIGITITICA AIACGAITTA AIAICGAICA GAGIAATICI ITHETACATT GAAATGAGG GCITIGGITTA AAAAAAGACC TITICCCICIC CCICCCCTA GAACAACCAG TATTAGAAGG TGCCACCATT GGTGCTGCCT TCTCTTCCCA CAGCCTGTAA CICAGIGITI IGIACTICAC IGAATIGIGA IGGITAGAAA CITICGIGGAI AGTITIGIGGA AATCATCCAA TTAAACATAC TGCTTAAAAC AGTOTTGCTG TGACTTCAGA GACAAGCCTG GOTOTITICOC ATGRECCECE TOTETACTEA CCAATCAGTG TOGCATGAGG CCCACGCCAC CCAAACCTIT CACTITICCAA AGAGCIAGCC GTCCTCCACC CAGTACCATG TCCTAGCCTG GAAGGGCAC CITAGGAAGC CCCTICGCTT CAGTIGCTCG CTICTGGGTG TGCTCCCTTC GAAGGCCCAG ATAAGACAGG GAACACTTGT GAGCACACAG AGCAGCATCT GATGCCCTGT TCTGCATTTG TTAGTGGTAA TATTCTTTAT GTATAATAAA TTTTTATACC CAAAAAAAA AAAAAAAAA ACTCGA

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1140 1200 1260 1320 1380 1440 1500 1516

## (2) INFORMATION FOR SEQ ID NO: 189:

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- (1) SEQUENCE CHARACTERISTICS:
  (A) LENGTH: 681 base pairs
  (B) TYPE: nucleic acid
  (C) STRANDEDNESS: double
  (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 189:

GCTCCCATGT TGCTGGCTGT CCGTACATCA CCCTGTCCCC TGCAGGAGGG GGCTACAGGC CATCTCCCTC CTGTAGGCCT CTGACTCCCC TCCACTTTTG GGCCCTCAGC TTATCTCGGG

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9 120

180 240 30 360 CAGGGGACCA TTGCAGCATC CTCCCTCCT CNGGACTCAA GGTGCTGAGG TATAAGCCCT GGGCCCCAGA TCCCTGRTKA CACCTTCCTG GAGAAGACTC TCAAAAGTGA CTGTATATTT GAGTICACCA GCAATAACTC CCCACACTCG AAGCAGGTCC AAACCCMAGG ATCCCAGGGT CCTTGGGCTC TGTGGCACTG TCTTCCCAAG ATCCTTCCTG TTGCACAATG GGAAACCTAA 8

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쏬 6 30 25 20 5 55 80 3 5 S TATTOGCTAC ACTICIDAAC AACCICAAAG TICAAGAAAG GCAGAACAGA GITTGTACCA AGCACATGTC ACTIGGGTT TATGGATTIG GTIGTGAAGA TICGCIGAAT CACTIGTICA TCTIGITIGA ATATATIGGT GAAAIGGGAA AAGACTACAT TTATGCCGTA ACACCGTTAC GCTGGATGAT TGCTGGGCAA AGGTGGCCTT TTAGAGCTCT TAAAAAGCCCA CAAAAAGGCT GITAGICTAC AAACCATIGC AGCAGCACCA ICAAACCAGA GICIGCCACI TITIGICAIC TOGTICTICCTC CTGGATGCTG CTCCTTGAAT YTTTTTTTTT GAWAAACCYT TTAMAATTAA CAGCITICCIG TOGGGIOCIG ACATOTOTCA CCACIOCCCC CCITICCCCC GGGGGGTCA TATOTTIACA TECECGAGGG GITTETGEET CETECCEACE CAGGICAGGG TOTOGTCCAG GTCTGTTTCA CCCAGCCCGG AAAGTCAGAG ATGTATATTG GAAAATTTAC AACTCCATCT CCCTAGAGGG CCTGAGAGTT GCTATTOGAC CATGTAGAAT GTTGCAATAT TGTTTACAGG TIGAAGAIGC TITAATOGAT AGAGACCITG TACACAGACA GACOGCIAGI GCAGIOGIAC TGAATGAATA CAGAGITCCT GAACTGAATG TICAAAATGG AGTGITAAAA TCGCITICCI CIGINGCAAT AGCIATIGIT GCAGANACAT GITCACCCIT TACAGIACIC CCIGCCITAA GCCTCAAGCC ACGCATATGA TAATTTTICTG GAACATTCAA ATTCAGTGTT TCTACAGCCA AAAAAAAAAA AAAAAACTCG A TOTOYOTOCO TOGOTOGAAT GTOGGCOOCT GCTOCOOGTO AGGITGTOCT GTOTOTOACO GAGGAAAAAG ACAGGGGCCT GCTTGCCCAG CCATGCGAGG GATTCCATGC CCACCTGCCC TIGCAGATCA GIGTAGGACI, GGICCATAGG GGAAGAGCTA GGAANTCCAT AGGC TAATGCACAS TACTICACAC CITAAACITG CITIGATITG GIGATGIAAA CITITAAACI racacctata ticgitatga actigactat atcitataat titatigitiw attikgigki ACTATOTATO GCCCAATOTR TTTGAGACAT CTCCTCATOT AATTCAGGCA GTTATGGGAG ATTOGTAGAG CCACAGTOAA CACATTTOGT TATATTOCAA AGGCCATTOG COTOATGATG 2 ACATTGGTTC CCAGGACGCT CTCATAGCAC ATTACCCAAG AATCTACCAA CGATGATAAG INFORMATION FOR SEQ ID NO: 190: Ξ Ĕ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 190: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 1014 base pairs 1014 960 900 840 780 720 660 600 540 480 420 360 300 240 180 189 60

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68 600 540 480 420 5 S INFORMATION FOR SEQ ID NO: 191: E

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2779 base pairs (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

25 20 5 CTOCTOTICAG COCCOGCAGO CCCTCCCCGGC TTCACTTCCT CCCGCAGCCC CTGCTACTGA GICCIDACIG ANCIDAGONA COTOCIOCIO CINAGANIGAO CICACOCIOS AGCIGIODAS GCCAACACCG ACGCGCANTG GGAGGAAGAC AGGACCCTTG ACATCTCCAT CTGCACAGAG GAAGCTCCGG GATCCCAGCA GCCGCCACGC CCTGGCCTCA GCCTGCGGGG CTCCAGTCAG CCAGAACCTO CTTGCTGGAG CTTAGTGCTC AGAGCTGGGG AGGGAGGTTC CGCCGCTCCT ASTINAGOTO COGVICTOCA COSTGCOGGO TGGCCAGGIG GGCTGAGGGT GACCGAGAGA TOGERGERGG GTGTGTCCRG ATGGTCRGTC TCTGGTGGCT AGCCTGTCCT GRCRGGGGRG SEQUENCE DESCRIPTION: SEQ ID NO: 420 360 300 240 180 120 8

8 S ş 3 6 35 မ GAGIOGIGCT AUGGGCCIOT CCGGGIGICG CIGIAIGACC IGGCIICIGI GGACAGCIGI GUAGAACCCA CACCAGCCCG CCAGCCTGCA GGCACTGACT CCCAGGGCAA CACAGTCCTG ACGOGOTCAA TOCCTOCATT CTGCCACTGC TGCAGATCGA CCGGGACTCT GGCAATCCTC CGCAACCTGC AGGATCTCAC GCCTCTGAAG CTGGCCGCCA AGGAGGGCAA GATCGAQATT CATIGOCCTAG TGATGATOTO GGACAACTCA GOTGAGAACA TIGOACTGGT GACCAGCATG AATACACAGA GOOCTCCACA GGTAAGACGT GCCTGATGAA GGCTGTGCTG AACCTTAAGG AGGATOTOGO TOGACTITOCA GAGTACOTGA GOAAGACOAG CAAGTACOTO ACOGACTOGO COGATICIAMA COGATITIGAE CGAGATICOGE TETTICAATIGE GGTETICECOG AATTOGCOCC TCAGATAAGA GTCAACCICA ACTACCGAAA GGGAACAGGT GCCAGTCAGC AGCIGGATITI TOGGAGCGGG CIOCCICCCA TOGAGICACA GIICCAGGGC GAGGACCGG ITICAGGEACA TECTIGEAGEG GGAGITITICA GGACTIGAGEC ACCITITECEG AAAGITICAEC TATGATGGGC TOCTOCAAGC TGGGGCCCGC CTCTGCCCTA CCGTGCAGCT TGAGGACATC IGAGCTACCC CICTYTTIGG CCGCTIGCAC CAAGCAGIGG GAIGIGGIAA GCIACCICCI IGCANGCCCO GONCTOCOGO GONTOTTOCCA GAAGGGOCCAA GGGACTIGON TITTATIVICOG Indicattiga gaagaggagw chocaghdig tgaagcheet gonggagaat goggeeaang AGCCCCTGGT AMATGCCCAG TGCACAGATG ACTATTACCG AGGCCACAGC GCTCTGCACA ITITICAGOTT GGAGACATTA GATGGAGGCC AAGAAGATGG CTCTGAGGCG GACAGAGGAA COLOLOGO 1440 1380 1200 1140 1080 1020 1320 1260 960 900 840 780 720 660 9 540 480

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8 120 180 240 300 360 420 480 540 9 99 720

1500 1560 1620 1680 1800 1860 1920 1980 2040 2100 2160 2220 2280 2340 2400 2460 2520 2580 2640 2700 2760 2779 GAGGAGAACT CAGTGCTGGA GATCATTGCC TTTCATTGCA AGAGCCCGCA CCGACACGGA ATGGTCGTTT TGGAGCCCCT GAACAAACTG CTGCAGGCGA AATGGGATCT GCTCATCCCC TACCATCAGC CTACCCTGAA GAAGCAGGCC GCCCTCACC TGAAAGCGGA GGTTGGAAAC AAGITICITOT TAAACITICCI GIGTAATCIG AICTACAIGI ICAICITICAC CGCIGITIGC TOCATIGOTIGO TGACGGGCCA CATOCITIATO CTGOTIAGOGG GGALCTIACOT COTOGNGGGC CAGCITCIOGI ACTICIGGOG GOGCCACGIG ITCAICIGGA ICICOTICAI AGACAGCIAC TITICAAAICC ICITICCIGIT CCARGCCCTG CICACAGIGG IGICCCARGI GCIGIGITIC CTGGSCATCG AGTGGTACCT GCCCTGCTT GTGTCTGCGC TGGTGCTGGG CTGGCTGAAC CTOCITIACY ATACACGIGG CITCCAGCAC ACAGGCATCY ACAGTOTCAT GATCCAGAAG CCTGAGCCAG GANNTTGGCG CCCCGAAGCT CCTACAGGCC CCAATGCCAC CAGCCCATGG AGGACAGGA KGAGGAKGGC NACGGGGCCC AGTACAGGGG GCCTCCTITGG AGCTCTTCAA ATTCACCATC GGCATGGGCG AGCTGGCCTT CTGCACTTCC GCGCCATGGT GCTGCTGCTG CTGCTGGSCT ACGTGCTGCT CACCTACATC CTGCTGCTCA ACATGCTCAT GGCCTCATG AGGGAGGG TCAACAGTGT COCCACTGAC AGCTGGAGCA TCTGGAAGCT GCAGAAAGCC ATCTCTGTCC TGGAGATGGA GAATGCCTAT TGGTGGTGCA GGAAGAAGCA GCGGGCAGGT GTGATGCTGA CCGTTGGCAC GGAAAACTAT GTGCCCGTCC AGCTCCTCCA GTCCAACTGA TGGCCCAGAT GCAGCAGGAG TAAGCCAGAT GGCAGCCCSG ATGAGCGCTG GTGCTTCAGG GTGGAGGAGS TGAACTGGGC ITCATOGGAG CAGACGCTOC CTACGCTOTG TGAGGACCCG TCAGGGGCAG GTGTCCCTCG AACTETECAA AACCETGICE IGGETTCCCE TEECAAGGAG GAIGAGGAIG GIGCETETGA OCCAGAGGAC AGAGCAGAGG ATCTTTCCAA CCACATCTGC TGGCTCTGGG GTCCCAGTGA AAAAAAAAA AAAAAGGC CCCTGGTGAG AGAGTCAGTG TATCCTGGAA CCAGGARCAG S

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INFORMATION FOR SEQ ID NO: 192; ĉ

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LENGTH: 1923 base pairs (A) LENGTH: 1913 base pair
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLGGY: linear SEQUENCE CHARACTERISTICS:  $\Xi$ 55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 192:

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TOPOLOGY: linear

1020 1080 1140 1200 1260 1320 1380 1440 1500 1560 1620 1680 1740 1800 GOGCTGCCCC TOGATCCTGC CCCTGCTCCT ACTCAGCGCC ATCGCCTTCG ACCOSCITCOS CITCOCCICOS CITORACOCOS CACOCOCOS CAACATGATIC COCTIGOGOCO CAGACGICCT AGAGCCTCAT GGAGTACGCG TGGGGTAGAG CAGCGGCTGC CATGCTCTTC TGTGGCTTCA TOSTAATITA CCCOSTGAAG TACACCCAGA CCTTCACCCT TCATGCCAAC CSTGCTGTCA GIGCCITICIT CITICIFICATIC CICCCCAACT ACGAAGATGA CCITICITGGGC AATGCCAAGC CAGATICCTTC TCTTCCTGAG AGTGATTGGA GGTCTCTTG CCTTGGCTGC TGTGTTCCAG ATCATCTCCC CCAGGTACTT CTACACATCT GCCTAACTTG GGAATGAATG TGGGAGAAAA TCGCTGCTGC CTANITIACCT ATACTATICC AATAITITICCT TATAICTATC CATAACATIT ATACTACATT IGAGATGGAC TCCAGAAGAA GAAACTGTTT CTCCAGGCGA CTTTGAACCC ATTTTTTGGC AGTOTICATA TTATTAAACT AGTCAAAAT GCTAAAATAA TTTGGGAGAA AATATTTTT AAGTAGTGIT ATAGITICAT GITTAICTIT TAITAIGITT TGIGAAGITG 1GICITTITCA TGTAAGAGAA TATGCACGTG AAACTTAACA CTTTATAAGG TAAAAATGAG GTTTCCAAGA ITTAATAATC IGAICAAGIT CITGITAITI CCAAATAGAA IGGACICGGI CIGITAAGGG CTAAGGAGAA GAGGAAGATA AGGTTAAAAG TTGTTAATGA CCAAACATTC TAAAAGAAAT AAATGCATAT CATTTOTGAG AATTTCTCAT TAATATCCTG AATCATTCAT TTCAGCTAAG STITATATGI ICAGAACCAG AGIAGACTGG AFTGAAAGAI GGACTGGGTC TAAITTAICA GCAAAAAAA AGTITATITI CAAGCCTICG AACTATITAA GGAAAGCAAA ATCATITICCI GCTTCATGTT GACTCGATAT GTCATCTAGG AAAGTACTAT TTCATGGTCC AAACCTGTTG TGACTGATAG ATCTGGTTAA GTTGTGTAGT AAAGCATTAG GAGGGTCATT CTTGTCACAA AAGTGCCACT AAAACAGCCT CAGGAGAATA AATGACTTGC TTTTCTAAAT CTCAGGTTTA ICTOGGETET ATCATATAGA CAGGETTETG ATAGITTICCA ACTGIAAGCA GAAACETACA TATAGITAAA AICCIGGICI TICITGGIAA ACAGAITITA AAIGICIGAI AIAAAAAGIG CCATAGITGG TAAGGCTITC CITTAAGIGT GAAATATITA GATGAAATIT TCTCTTTTAA CCACAGGAGA ATTCGGGGAT TTGAGTTTCT CTGAATAGCA TATATATGAT GCATCGGATA GGTCATTATG ATTITITACC ATTICGACTT ACATAATGAA AACCAATTCA TITIAAATAT agtictitat aggottaggg tgtgggaaa tgctatatta ataaatctgt agtgtyttgt GCTGGCCGC CGCGCTGGT TGCAGTCTAG CGACCACGGC CACTISTISSTIC GAAATICCTCC CAAGAGGGCG GCGGCAGCGG GTCCTACGAG CITACATCIA TAACTGGGCC TACGGCTTTG GGTGGGCAGC CACGATTATC TOTTOGCCCT CTGTGGACCC TCATCCTGGT GATCTGTTTC ATCCTCTCT TGGCCTGCGA ACATCATCCC Š 2 13 8 23 8 33 **\$** 5 S 25 8

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35 8 55 50 3 6 30 25 20 15 5 S CICCCCGGCC CCITITICGCC CCCTGCCTGC CATACTGCGC CTAACTCGGT ATTAATCCAA TITTICIAGCA TCCTCTTAAT GIGCAGCAAA AGCAGGCRAC AAAAICICCT GGCTTTACAG ACCGATITIT AAAGIIGGIG CAICTAGAAA GCTITGAAIG CAGAAGCAAA CAAGCTIGAT GENTICATOT CHARGETAGA GECATITIGA ACAACAANIC TACGIAGITA ACITGAAGA GGTAATAGTG ACATGCAGGC ACCYCTTTTA AACAGGCAAA ACAGGAAGGG GGAAAAGGTG AACTGAGGAG AAGCTGATCC AGTTTCCGGA AACAAAATCC TTTTCTCATT TGGGGAGGG ATCCTOGTTC AMACGGGTGC CTGGTCAGAA GGCCAGCCGC CCACTTCCCG TTTCCTCTTT GCCAMAMACA AGACGCGTAC AGCACACACT TCACAMAGCC AAGCCTAGGC CGCCCTGAGC CTCTTCCATT AACCAGTGGC CGGTTGCCAC TCTCCTCCCC TCCCTCAGAG ACACCAAACT AGCTTATTTT GTAAGAGTGA GCTCTGGTGG AGACAAATGA GGTCTATTAC GTGGGTGCCC CCTGCTAGGC GGCCTGCCCA GCTGCCGCCC CCGGACTCTG ATCTCTGTAG TGGCCCCCTC ACATCACCC CAACTCGGGT GCTGCTGGGA ACAGNGCCGG GCCCAAGTCC ATGGAGGTCT GCTCCCACAC CAGCGAGGGC ACCCGCAGCC GCTCGCACAC CAGCGAGGGG GCCCACCTGG AGOSTICATTO TOTOGATOGO ACCOGUAGOO GOTOCOACAO CAGOGAGOGO ACCOGAAGOO CCCCTAGTIGC CCTCGGCTAG CATGACCCGC CTGATGCGWT SCCGCACAGC CTCTGGTTCC COOCATISCET GECTICATICAL CIROCAGIET GECTIGECETG CITAACCOGET GICTETTOTI GCTGCAGGG ATCAAGTCTT CTCTGGGGCT GGGCACGTAN AAGAGCATGT GGCTGGTGGA GGCCTTGCCA AGGCTCAGGG GGACACTCTC AAAATTACAC AGCTTTTAAC AGGTGGCAGA ATTGGGGTTC Ą AGACCCAGAT CTGGGTTCAA GTCACTCATG GTGTGATTGC GGCATTCCTT CCCGCATCTG (2) INFORMATION FOR SEQ ID NO: 193: CAGATTATTA TITTOTAAGT TOTOGAAAAA GCTAATIGTA GITTICATTA TGAAGTITTC CCAATAAACC AGGTATTCTA AAAAAAAAAAAA AAAAAAAACTN GAGGGGGGG CCGGTACCCA ž E TCTCTCTCTC CGAGTGGACA TGGAGAGGAC GGGGGCCCAG CAGCTGGATG SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 193: (C) STRANDEDNESS: do: (B (A) LENGTH: 2346 base pairs STRANDEDNESS: double TYPE: nucleic acid 1200 1260 1080 1020 1923 1920 900 840 720 660 60 540 1860 780 480 420 360 300 240 180 120

> 8 35 30 25 20 ᇐ 10 S ACAATC GOCCTICCCT GOCCTICAGT AGAAAAGICG GOCATCGGGG CAAGAGAGGC TGAGTACGGA GGCGGAATTG GGGTTACTCG ATGTAAGGGA TICCTTGTTG TIGTGTTGAG ATCCAGTGCA AMAMMAMA AMAMMACT CHAGOGGGC CCGTACCCAA TICGCCGTAT AIGAICGTAA TITICCAGACC AATAAATITIG TAACITITIGCA AAAAAAAAA AAAAAAAAAA AAAAAAAAAAA TOCAGOCTTO CAGGTGGCGT GAGAGGCAAT GACTCGTTAC CTGCCGCCCA TCACCTTGGA GTANGTCAGC CACTGGGACC CGAGGATTTC TGGGACCCCG CAGTTGGGAG GAGGAAGTAG TICATITIGG CICACCOIGG ATITICICAT AGGAAGTIIG GICAGAGIGA ATIGAATATI CCTICCCICT CCIGCITCCT CITTICCIGC ICCCIAACCT TICGCGAAT GGGGCAGCAC TOGGAAACTA TTOTOCACAA GTCTTTCCAG AGGAGTTTCT TAATGAGATA TTTGTATTTA CACIGACGIT ICTOBBOGGO CAGIGOGGOT GCCAGGITICO IGTACIACIO COTIGIACIT COTOCOCACO GOTOCOTGAG AAAANOTAAG GATTOTGGAA TACATATICO ANGOGACTIT COGREACOTT CITOGARICCT CAGAACICIT TOCTICTICIC GOGGIGGOGG TOGGAACTCA ADTITICAAT CITCGACAGC TOGGCTGGAA COTGAACTCA GTAGCTGAAC CIGICTGACC CAGCTOCCCT OCCIOTISATG AGTOTCCTTG CAGGGGCCGG AGTAGGAGCA CTGGGGTGGG TTCAGGGAGA AGATCATTTA GATTTGTTTT GCATTCCTTA GAATGGAGGG CAACATTCCA TOCCTOTOCT COACAGOOCO AACOTOCOAC COCTGATACA TGAGOOAGTG ATTATTOTTIG OTTOTIGATIT CTOTIGGATICC CAGCITIGOTT CCAGGAATIT TOTIGTGATTG GCTTAAATICC ACAAAAATAT TTCAGCAAAC GTIGGGCATC ATGGTTTTTIG AAGGCTITAG TICTGCTTTC 2340 2280 2220 2160 2100 2040 1980 1920 1860 1800 1740 1680 1620 1560 1500 1440 1380 1320

(2) INFORMATION FOR SEQ ID NO: 194:

55 3 50 AGCTOMAGGT GCCCTGTGGN ACGAGCAACT GGACTATAGC AGGGCTGGGC TCTGTCTTCC TATCTGAACC ACCCTTTATT CTACATATGA TAGGCAGCAC TGAAATATCC TAACCCCCTA TOGTCATAGO CICACICITI CCCCCAAATC TICCICIOGA GCITTOCAGC CAAGGIGCTA Ξ (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 194; SEQUENCE CHARACTERISTICS: (A) LENGTH: 3054 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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AAAGGAATAG GTAGGAGACC TCTTCTATCT AATCCTTAAA AGCATAATGT TGAACATTCA

180 240

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	425			426	
	TICAACAGCT GATGCCCTAT AACCCTGCC TGGATTICTT CCTATTAGGC TATAAGAGT	300	J	GACADAGADA GADADAGATA TCADAGGCAG DADGGAGATC ATTINGTTOG GTCTGADAGG	2100
	AGCAAGATCT TTACATAATT CAGAGTGGTT TCATTGCCTT CCTACCCTCT CTAATGGCCC	360		NANAGICTITI GCINICCGAC ATGINCTGCT AGINCCTGIN AGCATTTTAG GICCCAGNAT	2160
5	CTCCAITTAT TIGACTAANG CATCACACA TGGCACTAGC ATTATACCAA GAGTATGAGA	420	8	GGAAAAAAA ATCAGCTATT GGTAATATAA TAATGTCCTT TCCCTGGAGT CAGTTTTTT	2220
	ANTACAGTEC TITATGCCTC TAACATTACT GCCTTCAGTA TCAAGGCTGC CTGGAGAAAG	480	~	AAAAAGTTAA CTCTTAGTTT TTACTTGTTT AATTCTAAAA GAGAAGGGAG CTGAGGCCAT	2280
9	GATGOCAGOC TCAGGGCTTC CTTATGTCCT CCACCACAAG AGCTCCTTGA TGAAGGTCAT	. 095	5	TCCCTGTAGG AGTAAAGATA AAAGGATAGG AAAAGATTCA AAGCTCTAAT AGAGTCACAG	2340
2	CITITICCCC TATCCTOTIC TICCCCTCCC CGCTCCTAAT GGTACGTGGG TACCCAGGCT	009	_	CTTTCCCAGG TATAAAACCT AAAATTAAGA AGTACAATAA GCAGAGGTGG AAAATGATCT	2400
	GOTTCTTGGG CTAGGTAGTG GGGACCAAGT TCATTACCTC CCTATCAGTT CTAGCATAGT	099	•	AGTICCIGAT AGCIACCCAC AGAGCAAGIG ATTIATAAAT TIGAAATCCA AACIACITIC	2460
15	AAACTACOGT ACCAGTOTTA GTOGGAAGAG CTOGGTTTTC CTAGTATACC CACTGCATOC	720	15 1	TRANTAICAC TITIGOTOTOC ATTITITOCCA GGACAGGAAA TATGITOCOCO COTAACITITO	2520
	TACTOCTACC TGGTCAACCC GCTGCTTCCA GGTATGGGAC CTGCTAAGTG TGGAATTACC	780		TTGCTTCAAA AATTAAAATC CAGCATCCCA AGATCATTCT ACAAGTAATT TTGCACAGAC	2580
ç	TCATANGGGA GAGGGAAATA CAAGGAGGC CTCTGGTGTT CCTGGCCTCA GCCAGCTGCC	840	<b>*</b> 02	ATCTCCTCAC CCCAGTGCCT GTCTGGAGCT CACCCAAGCA CTTGGTTGTG	2640
3	CACAAGCCAT AAACCAATAA AACAAGAATA CTGAGTCAGT TITTITATCTG GGTTCTCTTC	006	-	AACCIMACTG CCTTAACCTT CTGGGGGAGG GGGATTAGCT AGACTAGGAG ACCAGAAGTG	2700
	ATTCCCACTG CACTTGGTGC TGCTTTGGCT GACTGGGAAC ACCCCATAAC TACAGAGTCT	096	•	AATGGGAAAG GGTGAGGACT TCACAATGTT GGCCTGTCAG AGCTTGATTA GAAGCCAAGA	2760
25	GACAGGAAGA CTGGAGACTG/TCCACTTCTA GCTCGGAACT TACTGTGTAA ATAAACTTTC	1020	25 c	CHOTGGCHGC AAAGGAAGAC TTGGCCCAGG AAAAACCTGT GGGTTGTGCT AATTTCTGTC	2820
	AGAACTGCTA CCATGAAATG AAAATGCCAC ATTITGCTIT ATAATTTCTA COCATGTIGG	1080	Ü	CAGADARIAG GGTGGACAGA AGCTTGTGGG GTGCATGGAG GAATTGGGAC CTGGTTAIGT	2880
Ş	GAMANACTIGG CTTTTTCCCA GCCCTTTCCA GGGCATAAAA CTCAACCCCT TCGATAGCAA	1140	T.	TOTTAITCIC GEACTGIGAA TITIGGIGAT GIAAAACAGA ATAITCIGTA AACCTAAIGT	2940
3	GTCCCATCAG CCTATTATIT TITTAAAGAA AACTTGCACT TGTTTTTCTT TITACAGTTA	1200	_	ctgtataaat aatgagggtt aacacagtaa aatattcaat aagaagtcaa aadaaaaaa	3000
	CTTCCTTCCT GCCCCAAAAT TATAAACTCT AAGTGTAAAA AAAAGTCTTA ACAACACTT	1360	4	AAAAACTOG AGGGGGGCC CGGTACCCAA TITINCCAAAT AGAGATNSTA TTAC	3054
35	CTTGCTTGTA AAAATATGTA TTATACATCT GTATTTTTAA ATTCTGCTCC TGAAAAATGA	1320	35		
	CTGTCCCATT CTCCACTCAC TGCATTTGGG GCCTTTCCCA TTGGTCTGCA TGTCTTTTAT	1380	_	2) INPORNATION FOR SED ID ND: 195:	
40	CATTOCAGOC CAGTGGACAG AGGGAGAAGG GAGAACAGGG GTCGCCAACA CTTGTGTTGC	1440	. 64	(1) CENTENNY CHADACHTC.	
}	TITCTGACTG ATCCTGAACA AGAAAGAGTA ACACTGAGGC GCTCGCCTCCC ATGCACACT	1500	?		
	CTCCAAAACA CTTATCCTCC TGCAAGAGTG GGCTTTCCAG GGTCTTTACT GGGAAGCAGT	1560		-	
45	TAAGOCCCCT CCTCACCCCT TCCTTTTTC TTTCTTTACT CCTTTGGCTT CAAAGGATTT	1620	45		
	TOGADAGAA ACAITATOCT TTACACTCAT TTTCAATTTC TAAATTTOCA GOGGATACTO	1680	в	GECAGAGCTC GTGGCCGANA CTTTTTCTGC TCCTGGCTGC CACCTACTGG CTGGCCGGG	09
20	AAAAITAGGG CAGGTGGCCT AAGGCTGCTG TAAAGTTGAG GGGAGAGGAA ATCTTAAGAT	1740	50 c	CCCTOGCCTO GGCTGCACC AGCTGCGAG CGGGCTCCCA CAGCAGCCCC CTTCCAAGCA	120
	TACAAGATAA AAACGAATC CCCTAAACAA AAAGAACAAT AGAACTGGTC TTCCATTTTG	1800	o	GGTCCCCAC ACCGCGCACC TTCTGCGGGA ACGTGCTCGC CGTGCCGGGG ACCATATGGA	180
	CCACCTITICC TGTTCATGAC AGCTACTAAC CTGGAGACAG TAACATTTCA TTAACCAAAG	1860		CSGAAGSCTT TOTGCTCACC TACAAGCTGG STGAGCAGGS TGCCAAGCAGC CTGTTGATCC	240
22	AAAGTOGOTIC ACCIGACCTIC TGAAGAGCTIG AGTACTICAGG CCACTICCAAT CACCCTACAA	1920	55 <sub>T</sub>	TETTOSCHEC TOCTOGRACA COMOCOSOST TICTOCTOCC CANOTICOGRAC TOTOGRAPHICS	300
	GATOCCAAGG AGSTCCCAGG AAGTCCAGGT CCTTAAACTG AGGCTAGACA ATAAACCTGG	1980	F	חסוספחופרו פוספורוערו בפאוספרוספ בתכבוסכבום פפוסספאבבר חסבונסברא	360
9	GCAAGTIGAGG CAAGAGAAT GAGGAAGAAT CCATCTOTGA GOTGACAGGC AAGGATGAAA	2040	° 09	GOACTIGGIAIA CTIGCTIGCCTC TOTIGLIGOTICG GTIGCTIGCGCT TCCGCCTCGG 0000CTAGCC	420

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8 35 25 20 S 3 30 15 5 S TICADOGAGA TOGOCAAGOG GICCCOTOGO GOCTOTOGOCA GCGGGCTTAT COTTOTOTO CCTCTGTGGT CCACCAGCGT CTTGGCTTGG TGGGAGGGCT CTCAATCAGC AGGGCCCCAG CTYTIGGSCGG AGGGCCTIGGC TGATGGGTTG GGGCCACATC CCTGCTTCTT GCTCCTGCTC TTAACACCAG CCCTCCAGCA TCTAATAGAC TTGAATCTAC TCTAAACGAA TATTTAATCC TOTOGRAGOCA OGCOTOCTOC AGGCOATGOT COTTGRAGOT GGARATGTOR ACCOGRAGOCO AACAATO CTOTOCATAC GTGTGAGTCC AGCTAAAAAAG ACAAAAACAGA ACCCGTGGGC CCAGCTCGG AACCTCAACT ACATTGTAGC TCAGTCCAAC GACTAACCCT GAAATGGGGG TGTTCCAGCC KAGGGCAAGA AGAAGTOGGG CAAAGCCTOG CGCTCOGCCG CGGTCGCGGC AGCTTTGCAA GOCACGAGGA OCGACAGGGA GTGGGCAAGG GGAAGAAGCA GCTTATTTGA CTAACCAGCC AAAAAAAAA AAAAAAACTG GAGGGGGGC CCGGTACCCA AATCGCCGGA TATGATCGTI ANCONONIO CONTROCCON NOVALLO GACCIAGOAC COAGCACONT NOVALGO AGGCCACACA CTACAGCCTT CTGGCCACGC TGGAGCTGCT GGGGAAGCTG CTGCTGGGCA CACCACAGIC ACCITICACIO GGAATGATOC GCTGCAGCCA GCTGGCCCCC AGGGCCTTGC AATCTIGAGA GOGICAGCCT IGCIGAGCCI AIGICIGCAG CACTICTIGG GARGCCIGGI AGGTGCGTGG AGAAGGCTCC GACGTCTCCG AAGTGCAGCC CTTGGGATGG CATTCCGTTC TOCCAACCIT GCCGTCCGAC CTCCTCCGCC CCCATGCGGT GACCCCGTCC GTGTCTGTGT (2) INFORMATION FOR SEQ ID NO: 196: TOTCAGACTO CCTTGGTCTT CCACCTTGGA CACCCTGGGG GCCAGCATGG ACGCTGGGAC E X. SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 196: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double g (A) LENGTH: 1290 base pairs TOPOLOGY: linear 540 480 420 360 300 240 180 120 600 480

8 2 INFORMATION FOR SEQ ID NO: 197:

15

AGGGGANCCC CCCATTIAAA ATTITNGGIN

CCIAMCIATT TITITIOGOGG GICCCINAAG GICCCCCTAA AANCITITIT COGAACCCINA

1260 1290

1200 1140 1080 1020

AAAGGGGAAA AAACSYTTTT YTGGGGGGGAA ANGGGGGCCCC CNTACTTTNA ACAYCCCCCC

5

TICCCCAYIT TIGGAAACAA AWIYCCCCCT TITIAAAAAA GIIGGAACCC CCAMCCYTCC

COCCADATICE CEYTTATTAA TICCAAAAAA ATAAACESAA AAWOGGITIG AATTITITIKT

TOCKOGITWA AWIITIIGIT TAAAICARCI CAATITITIT AACCCAATAA GSCCGAAATC

TTTGCGCAGC CCTGAATGGC GAAATGGCAA ATTGTAAGCG TTTAATATTT TKKTTAAAAT

960

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SEQUENCE CHARACTERISTICS (A) LENGTH: 1020 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: doubl.
(D) TOPOLOGY: linear STRANDEDNESS: double

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ၓ GOTOTOCCTG GATOGTCOTG TAGGTGAGTT TTACCAAGGA TTATGGTAAC AAATGAGTGA GACCTUTATO GAGAAAATAT TGAAGNINCAT TAAAGAAGAC CTCATANTAG GAGAGAATOT Ě SEQUENCE DESCRIPTION: SEQ ID NO: 197: 120 8

ATTATOTAGG TAGAAAAAA TOCAAGCAAG CTOTTAAAGA TOTTOGATOO CATTATATAG SCITTIGGAGO ATTIGIATIG AGCITTITACA GIATICATIT TICAACICAA GGCAATGGCI THIGIATAGC TGAAATCIGT AATTCAATCA CITTTICICT TITATCCTCT AACCAAAAA AUGCITAGIG IGIATITICIC ICITIGAGAC ACIGIAATIT CIACCAGAAA ITICCAGAGC TICTACACCA ACTICTAATICE ATAAACGGGT CITATGACAT CTATGAAGTA GTAGCAAGAC 240 180 420 360 300

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TOCTOATOAT TECCTITITT GREGITETAA ATAGACITAC TIGGACITTE AAGATGAGIT ACTOCTTOTO ATOTTACAAA TATOTOATAT GOTAATTTTO ATAACAGATG TOAOTTTTGA TIGITIAAIT TIGCATCCCA AATGITTITA AICTITGIAT AITTITIAAA AAYCCITTIC 600 540 480

ACCAAGAATT GGTGATTTOT TTATAAGAAA AAAACTGGCT TCATTTCTGT GAAATTGCTC TTIGAAAATT TCTTTTTACA COTOTAAGCC AACTGAGATA CCOTGATGOT OTTGATTTCT 660

720

780

GAACTITITIG GATATIGATA TIGATITIGAA AATATITITIGG AATTITITICT ACTIGAAATT TICAAIGAIG CITACCAICI ATITIAGCCA CIGAGCCITI TATTATITOT CTATTITOTAA AGTITÄTTIG TCTTAACICA TITAATAAAT ATACTGTTTA TCIGITTCIG AATGGOGACI

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TGTGCCTTAT TCÇTGGAGAA TCTGTATACG GCTCGCCTAT AAGAAATATA GCCTCTTCAI

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COTEOTOMAC TOGOMANACE CTOGOGTTTA CCCAACTTAA TOGOCTTOGA GCACATOCCC GOTACCCAAT TCGCCCAATA GIGAGICGTA TTACAATICA CIGGGCCGIC STITTAACAA 

840 780 720 660

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TTAGAAATCT AATKGAAAAT TCTATAATGT ACTGAAAGTA WGGTTGTGTA CAGTGAKCAC

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CTITICOCCAG CIGOCOTIAA TAGCGAAAAA NGCCCGCACC CGAATCGCCC TICCCAACAG

		PCT/US98/04493	WO 98/39448	PCT/US98/04493
	429		430	
	TCTCTAATAA TAITGATGACT TGCCCTAAAN GAGGAGGGAC ATGTCCCACT TTCCACCAGG	1020		
			(2) INFORMATION FOR SEQ ID NO: 200:	
8			5 (i) SEQUENCE CHARACTERISTICS:	
	(2) INPORMATION FOR SEQ ID NO: 198:			
. 5	ovančas (T)		(c) STRANDELNESS: GOUDLO (D) TOPOLOGY: linear	·
2	(A)		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 200:	
	(C) STRANDEARESS: double (D) TOPOLOGY: linear		CCAGGGAAGC CCCARGCCTG TCCTGAATTG ACATCAGTGC TTCCCTGAAC TGCCTCCCCC	09
15	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 198:	1	15 ассетивае аттагессая валасттать тттетьсял остальнае тестовеле	لَّتِد أَ 120
	ANTICOCGAA GCTGAGGGTT GTGTGCCATC GGGGAGCCA AGTCTTTTGA CCGGACCCTT	09	CAGGGACTGG TGCAGGTAGG CTCAGTCCT AGAAGGTCTC TGAAGATCTG	180
5	CCCCCCCCA AMANCTCAA GITGATITTGA GAGCCTOTKT TIGGGGTTRA GCCGAGCTGC	120	GACTGAGGAC CYTGCTACTC CCCAAGCCAG AGCCCATCAG CCAGGCCTGC TGTGAGCCAC	ac 240
3	TOCOCCTIY OTCCCCOSCC AGGACACAAG YTACTITICAA COCCOCCCC CCTCCTTAT	180	CTGCCTGTGG AGTGCTGAGC TCAACCAAAG GCTGGCAAGC TCTGGGCCTC ATTTAAGGGA	300
	GATISTICCTIC AACCLAGGGG CGGCCTCTIGC CCTCTACTICG TGCCAGGCCC ACTTGCCAGG	240	TICTICATIGAG COGATIGAGC CTGGAGGCAG COCATTAAAG CATCTGGCTC GTTTTTGGAA	AA 360
25	CAGGAGCCT CCCCAAGCCT TCAGGCTGC TCGGAGTCAC CTGTTGGAAT GGACTAANG	300	25 лалалалала далаль	376
	GACCETTOTG TOGGAACAGG TGCTCCAAAC ACCCTGCTGC TGGCTGCCAG GCAGGCCCTC	360		2
ç	TOGANGGOAA GOGGCAGGAC TCATCAGGAC CTCCTGGAC CCTGCAGGGC AGGCAGTTGG	420	30 13) TATENDAMMETAN END EDG TO NO. OO.	<del>,</del> .
ક	CCCGAGCCCA AGCATTTGGC TCTGCTTGCC CCAAGGGGAC AGGAAGCCTC TTGGGCCTCT	480	TO THE OWNER	
	TOCCTTOCTS GACAAGGCC CCTGCCTTTG CCTCACATA' ACTG	524		
35		3.	35 (c) STRANDENESS: double (D) TOPOLOGY: linear	
	(2) INPORNATION POR SEQ ID NO: 199:		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 201:	
40	(i) SEQUENCE CHARACTERISTICS:	40	O CCCAGTATAT TYCTATAACA TYTATYTYAG TGAACTTATA ATGTYTCTTT GPATTAAAUT	09
			ATTAGALTAL ATCITTAGAL AATALTOSTA CINDASTRAGT AGGIAATATA TAITTTAGTG	120
!	(C) STRANDEZNESS: double (D) TOPOLOGY: linear		AAAAATAAAT TOTGCATCTA ATGTCTACCA ATTAATGTAC TTGTAGATGT ATCTTATCT	180
45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 199:	ŕ	AACTICAGTC TITIGCTGCCC CTAATGAGGT GTGAAGGACT CTTCTCCCCT GGGGAAGTTT	T 240
	GTGATACAAG GAAGGSTGAT CATCATGT CACCATGCAA TTCCTGCTCA CAGCCTTTCT	. 09	TICITITINCA GGAGGAAGGA GGCCTTICCC AGGTAANGIG ICINGAGIGT TGGGCAGAAA	300
20	OTHOGHECA CTICHGCTC THISTGATOT COCATATOC CHAGGETTCT COCCCHOCTA	120 50	O ANICIOGGAC CACACCACAC CAGITICICIC CITAATCCAC GICATITICCC ITCINICCO	. g 360 <b>4</b> -
	GAAGGGCTTC TTGATAGATT AGAAATAAG AATGAGTGAC ATTTCCTATG TCCATATAAG	180 .	GCTANGITIC CAGIGICCIC IGGGIGITIC CAAGAGCAAC AAGAAAYGAA TAAAICICTG	rg 420
ď	AAGGAGCCAC AAGACATGTC TITTAAATAA AAGGACAGTG TCCATCCTTT TAACTGCCGA	240	KIGAGITGIT IAITIGITCI ICACITIGIT ITACACIGIA WITICIGAGI TIAIGGGIG	480
r r	ATAGAACCTT GOTCTCATCC TCCTGGAACT AGGSCTTAAA ACAGCTTCTG TGTTTCTSAT	300	CTGTGAATTA AAAAGGAAAA GTRGAAATAA GTAAAACTCA GGTTGAAGGA AATATACATA	PA 540
	TKGTCTCART GTTTTGCCAA GGTTTTATTC GG		AATAAGATAA AGCTGACCTG TAGATATARR CAGGTTATAA RAGCTTAGAG TTGTCTAAGT	ar 600
9		09	D TGRGTGCAAA KITICCICTG ATCITICTGA TGCCGARACA AAAAAGGCAG TCAITGTTTGT	99

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50 ટ 6 35 30 25 20 COGOGAGICA GGCAGTICCG TCACCAGCIA CICGAICIGA GGCACITICI AGIGIGITAG GTCAGTCCCC AAAGATOGAG AGCTTGAGTT CTCACAGAAT TGATGAAGAT GGAGAAAACA GITISCAAGOG CAGAGAAGAA ACGGTAGCAG AAGATGTITIG TAITIGATCTC ACTTGTGATT ATGACAAAAC AAAGGGAGAT GATACAGACA COMGGGATGA CATTAGTATT TTAGCCACTG AAAATGATAG TATCCTGATG AATCCAGCAC AGGATGGTGA AGTACAACTG AGTCAGAATG CACAGATTGA GGATACGGAA CCCATGTCTC CAGTTCTCAA TICTAAATTT GTTCCTGCTG ATATOGAGAG TOTTCCOTTG CACCITTCIC TGACTGAAAC TCAGTCCCA AGGTGGAAGA ANTCCCTGAG ACACCTTGTG AAAGTCAAGG AGAGGAACTC AAAGAAGAA ATCAGGAGGA AGCTATOGAA ATTAAAGAAC ACCATCCAGA GGAGGGGTCT TCAGGGTCTO ATCTTGGGCT ATCTTTGACA GGGGATTCTT GCAAGTTGAT GCTTTCTACA AGTGAATATA (2) INFORMATION FOR SEQ ID NO: 202: CCITATITCT TCTAAACICA CCATTAATCT GAATAATAGT CAAATTIAGG GG Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 202: (A) LENGTH: 589 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 589 540 480 420 360 300 240 180 120 6

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(2) INFORMATION FOR SEQ ID NO: 204:

AAAAAAA

AGGACACCAT TITTICCAGAG CTOCAGAGAG CACCTOGTOG GGAGGAAGAA GTOTAACTCA CCAGCCTCTG CTCTTATCTT TOTAATAAAT GTTAAAGCCA GAAAAAAAAA AAAAAAAAA

840 847 660 660 720 780

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TEACCOCCAC TETICATRACE CTCATEANAA ACACTETICAE TATICETICETA TIGGACGACCET CCAGCTETICA GITACAAGTIG CAGGCGACTIG GAGGCAGGAC TETITGGGTCC CTGGGAAAGA GGGTACTAGA GGTACCGCTG GCGGAGCTGA

AGAACTOGGT ATGAGGCTGG GGCGGGGCTG GAGGTGGCGC CCCCTGGTGG GACAACAAAG

20

GESGACECTE ECTAMAGTES GEAGECCTTE GCCCACCTCA GECCCAGGT GEGAACATEG

CANGIOCANG GACCAAAGOG GOCCIGOCIT GCAIGGGIIG GCIIGCIGAT GGCIGCIGGA

5

TOCAGAACOC AGOCAGCAGT COAAGTAGCT GGACCCACGA GGAGGAACCA GGCTACTTTC

TOCCOGGGAA GCTGGATGCC CTGACTGAGC TGCTTAGCAC TGCCCTGGGG CCGAGCAGCT

CCCAGTACTG AGTOGTIGGAC ATCGTCTCTG CCACTCCTGA CCAGCCTGAA CAAAGCACCT

5

ATGACCTOCA GCAGAATCTO AGCAGCTCAC ACCGGGCCCT GGAGAAAACAG ATTGACACGC

GOCACGAGEG CAAGETGETG GEEGECATEA ACGEGTTEEG ECAGGTGEGG ETGAAACAEC GGAAGETEEG GGAACAAGTG AACTECATEG TGGACATETE CAAGATGEAC ATGATECTGT

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 203:

(C) STRANDEDNESS: double (D) TOPOLOGY: linear

CGTAATGTAT GCTCCTTACT CAAAGAGTGT GGTCCCAAAC AGCCTTTGGG AGGTCCTCCT AGTCATATTT TCTGGGAAAT ATTTCCAGTG TTTATTTGCA CTTTAGCCCA CTCTGTGTAG ACTOTOCACA COTTITIOTI AGITTATOTO TACATIGOAGG GIGIGOAGCA GOOTIGOAA TGATTCATGG ATGAAACCTG GAACATCTTO AGGACTGAGT TAACCATAGG TCCTTAAATA CTANAGENEG GEGTGTTGTG CAGEGGAMAT GGTCATETGE TGETAAAACA CAGETTECAT TICACACCOG CCIOGCAGIA AACACITOTA GIOTIGIGCA GIOGAAACOG ICATCIIICO TOYGTGCACC AAGTCTKAAC CACCACCTTC ATGGGACATA GRTTATOTGC TGGAACATAT WATGTGATTG GAATGGAACC CGARAAGAGA GCAYGCTGTG TTCTTGGGGA CAGGAAAGCT 1192 1140 1080 1020 960 900 840 780

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8 5 8 S 50 AGCTCACACC GGGCCCTGGA GAAACAGATT GACACGCTGG TOCHTOGTOG ACATOTOCHA GATOCHCATG ATOCTOTATG ACCTGCAGCA GAATOTGAGC GCCATCAACG CGTTCCGCCA GGTGCGGCTG AAACACCGGA AGCTCCGGGA ACAAGTGAAC ACAAACATAC TOOCAGGAAG GAGTCTCATG CTGCCCGCAG CATCAGCGCA ACNCNTGGCC AAGTAGCTOG ACCCACGNAG GAGGAACCAG GCTACTTTCC CCAGTACTGA GGTOGTOGAC ACTORACTOR TERGERETOR OCTOROGOROS AGGRAGOTES CRAGARICORAS CORACTAGINOS (xi) SEQUENCE DESCRIPTION: SEQ ID NO: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 852 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear CGGGGAAGCT GGATGCCCTG 180 360 90 240 120 8

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(2) INFORMATION FOR SEQ ID NO: 203:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 847 base pairs (B) TYPE: nucleic acid

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	433		434		
	ATRICOTOTOT TOCCACTOCIN TRIACCOAGO COTGAACAAA GCACCTCAAG TGCAAGGACO	420	AGSCETTGG CEACHTGAG ECCEAGETG GAACATGGTC ACCCCANTE TECAPARCT	096	
	AAAGGGGGCC CTGGCTTGGA GTGGGTTGGC TTGCTGAITGG CTGCTGGAGG GGAGGCTGGC	480	THE TAXABLE PROPERTY OF THE PR		
S	TAAAGTGGGK AGGCCTTGGC CCACCTGAGG CCCCAGGTGG GAACATGGTC ACCCCCACTC	240	CATCAMANAC ACTICACIA TACTACIATIO GACGACCICE AGENCICADI TACAAGISCA Socratementa constituente emperatorem procasamente custamatorem	<del></del> .	
	TOCHTACOCT CATCHAAAAC ACTICTCACTA TOCTGCTATG GAGGACCTCC AGCTCTCAGT	009	TROCETCE CONTRABO PARTICIONAL	3,000	
9	TACAAGTGCA GOGGACTGGA GGCAGGACTC CTGGGTCCCT GGGAAAGAGG GTACTAGGGG	. 099	GETGGCCC CCTGGTGGGA		
2	CCCGGATCCA GGATTCTGGG AGGCTTCAGT TACCGCTGGC CGAGCTGAAG AACTGGGTAT	720			
	сысствава свазаствая встватавая сысымама высысытт	780			
15	TTCCHGAGCT GCAGAGAGCA CCTGGTGGGG AGGAGAAGT GTAACTCACC AGCCTCTGCT	940	15 ACACTORANG TOTOMBARA BARACTORIA ANIAN		
	CTTATCTTTG TA	852		7	
20			20 C. TANDORMATION DOE GED IT NO. 206.		
	(2) INFORMATION FOR SEQ ID NO: 205:		100 Oct 17 700 Oct 100 To 100		
25	(1) SPQUENCE CHARACTERISTICS: (A) LENGTH: 1154 base pairs (B) TYPE: nucleic ecid		(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1178 base pairs (B) TYPE: nucleic acid (C) STRANDEINESS: double		
			(D) TOPOLOGY: linear		
9			(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 206:	-	
3			TCCCCAGGIG CACAGCCAGG GCCCTCCTGT CTGCAGGAGA AITCACAGCT GGTGTGGGAC	AC 60	
	GATTCOCCAC GAGGCTTGCT GGAGCAGGAG AAGTCTCTRG CCGGCTGGGC ACTGGTGCTG	09	TCAGCCCCTA GNCCATTCAA AGCCTAATG TTGTAATCAT ATCTTAGGTG TTGAAGACCT	CT: 120	
35	-		35 GACTOGAGAA ACAAAATOTO CAATAAGGYG AATTITIATCT TAGAGATCTG TGCAGCCTAT	AT 180	
	TOOGENSTEA ATSOCACTOG GCACCTTTCA GACACACTT GOCTGATCCC CATCACATTC	180	ITCTSTCACA AAAGITATAI TSTCIAATAA GAGAAGTCTI AATGGCCTCT STGAATAATG	TG 240	
Ş			TAACTICCAGT TACACGGTGA CTTTTAATAG CATACAGTGA TTTGATGAAA GGAGGTCAAA	300	
€	•		+U CAATGIGGGG AIGTOGIGGA AAGTIATOIT TOCCGCIOIT IGCIGIGGIC AITIGIGICIT	360	
	CIOSACITTA ACANGCAGA GAACCACOTG CACAACTICA TGATGGATAT CCAGTATACC	360	GCAGAAAGGA TGGCCCTGAT GCAGCAGCAG CGCCAGCTGT ANATAAAAA TAATTCACAC	AC 420	
45	AAGAGATGA AGGAGTCCGC TCCCCGAGTG CTACAAGAAG CCTGGATGTT CTACAAACAT		45 татсысысты ссамовсыст асметтевам масассысые мамсамасы атссымосст	480	
	ACTOGRAGIA AGRAGICTOA TOCTOCOCOC AGGCATCAGO GCAANCTIGOT GOCOGOCATO	480	ITCATCITAC AGGIGAACAA ACTOTGATGA TOCACATGTA TOTGITTTTGT AAGCTOTGAG	AG 540	
S			CACCETAACA AAATGTAAAT TTECCATTAT TAGGAAGTEC TEGTEGCAGT GAAGAAGCAC	AC 600	
2			OU CCAGGCCACT TGACTCCCAG TCTGGTGCCC TGTCTACACC AGACAACACA GGAGCTGGGT	. 099 TS	
	CACCGGGCCC TGGAGAACA GATTGACAGG CTGGCGGGGA AGCTGGATGC CCTGACTGAG	999	CAGATTCCCC TCAGCTGCTT AACAAGTTC CTCGAACAGA AAGTGCTTAC AAAGCTGCCT	720	
55	CTGCTTAGCA CTGCCTGGG GCCGAGGCAG CTTCCAGAAC CCAGCCAGCA GTCCAAGTAG	720	55 TCTCGGATAC TGAAAGGTCG AGTITTCTGA ACTGCACTGA TTTTATTGCA GTTGAAAAAA	780	
	CTOGALCCCAC GAGGAGGAAC CAGOCTACTT TCCCCAGTAC TGAGGTGGTG GACATCGTCT	780	AAAAAAACT ATTOCAAAGA TTTOAAGTIG TICTIGAGAGA TCTUCTICATIG		
	CTOCCACTOC TGANCCCAGC CCTGAACAAA GCACCTCAAG TGCAAGGACC AAAGGGGGCC	. 840	CIGACACACA ARCHITITAC TITIANICATA APPICATIVITY CICAACCAAT AAACTICAACA		
9	CTOSCITICA GIGGILIGGE TITOCIGATEG CTOCITICAGO GACOCITICO TAAAGIGGGK	006	09		

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35 ઝ 25 20 5 3 8 5 8 55 50 S CTGAGTTAGA ANTICACAAG TICTCCAGGT GATCTCATAC ATGCTAAAGT TIGAGAACCA AAAAACTOTO GACTITIOCCA TATAAGGGCT GTOGTTOTOT GTOGTCCCCT GGATIAAGAGG GAAAGITACA ATTOGAAAGT TICCIGCCAG CTICGGGAAT GACACTGCAA AGCTGATGCC GITIGCITITICA GITGCCTITTIA TITIGATITCCI GGAGAGAGCA GACTICGCACS AACATITCAAC CAGTETTEAG CTCNAGGGTT TTAAAA TIGGGCTTTT AAGGTTCAGA GACTGTGGGC TIGGGCACCT GCGCCCAGGG STTTTTGTGGG GITTICCCIGI TAACAAGAAA GICAGAGGIC AGTIGATCAG ACATTAGATI ATTIATIGCI TIGAGIAAAG TIAAIGCATT AAGAAGAGAI TAGATAGGGA IGGIGGCGTA ICTICCTACA TACTICTAGG ATTATAAGGA ATTAACATIG AGAIGACATT TCCATTIGAG AAGGAAAATA (2) INFORMATION FOR SEQ ID NO: 208: GGCCTTTGCC CCTTAGRAAA GTAGCTTTTA GGGGCAAAGA TTTGTTGATT TTCCCCCATTA TOCOGGAGE GTTCAGICTE GACCOTAGIC ACTGATTITE TOTAGITGIT AATAGAGIGG COTGAOTEAC CTCTCTATAG TOGGCOTGGC CGAGGCCGGG GTGACCCTGC CGAAGCCTCC (2) INFORMATION FOR SEQ ID NO: 209: AAAACTAAAA AAAATTAAAA AAAACTGGAG GGGGGCC CATCACCATT ATCTOGNAAC ATGCAGTAAA TGCAGATTNT TCATCTTCTC CCCAGACCTC ATTAACTAGT GCAGTTGACT AATTCTCTTT ACCTTTATCA TTTARGGTGA RGCATTGCAC AGAAACTOCC AGROTAATTC TOCTOATTAC TOCTOTACOC ACCCACTITC AGCTOCCCAA COCAGOGOTO ATATOACAGT AATOOTOAGA GGCAGAGOCCO AGCAGAAAAC AGCAATGOTA E (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 208: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 209: SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: (A) LENGTH: 697 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 932 base pairs
(B) TYPE: nucleic acid
(C) STRANDECNESS: double
(D) TOPOLOGY: linear 1166 1140 1080 1020 660 60 540 480 420 360 300 240 180 120 8

TCAGAATAAG TTACACACAA TOGCCACAGC AGTTTGTCTT TAATAGTATA GIGCCIATAC адараадаа аладарараа адарарараа, адараалаа адараласаа калалкаа AACAGCACCT TTTAATATAT AGGTCTCTCT GGAAGAGACC TAAATTAGAA AGAGAAAACT 1320 1260 1200 1140 1080 1020 960

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CATGAGACAG GATTATAGTG CCTTAACCGA TATATTTTGT GACTTAAAAA ATACATTTAA

AACTOCICIT CIGCICIAGI ACCAIGCITA GIGCAAAIGA TIAITICIAI GIACAACIGA

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(2) INFORMATION FOR SEQ ID NO: 207:

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Ξ SEQUENCE CHARACTERISTICS:

(D) TOPOLOGY: linear

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(A) LENCTH: 1166 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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GTAAGATTTT CCACACTACA GCTGGGTGTT TCTCTTTTCT AAAGTGAGGC CAGTGTTATT

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CCTTGTCACC ATCAGGCCTT TCTGGCTCCT GATAGGGTGG AGCAAAAGTG GAAAGGAAAA GIGSCCIGGG AAGGIGICCA CAGIGAGCCC IGIGIGCAGG ACTGICCACN ACGGITICACA

GAAAGAGGCY TITTICTCACA GCCATTATAT TAAATAGTAG GTCGATTCAC ATCYTCGTGC TOOTOGOOAC COTOCOOTOT GOOTOAGTGA CATGTAGATG ACTGAOTGCO AATAOTTGTO ACCATTCCCT GGAAGCAGCT ACCTAGGGGA AACAAGATOT AGTOCTATTG CCGATAACAA

3

TOCCAMEACE ATGEOCOGTO CTTAGGAAAT GAAAGAAGTE CEGGGTETGT CTETETCACT TGATTGTGGT CTAATTTCCA ACCIGCTCTG TTTTCTGTGA CATCTTGGAG GGGGAGCTAG GGGGCCCAGA GGCCGCCTTT TGAAATGTTT GCCTGTCTGA ACTGTGAAGA CACTTGGGAG GCTCAGTGGC TGAACAGCAT TOOCACAGCC TGCAAGTGTG TGTGTGTGTG AAAGAGAGAG AUGCIGCIGA CCIGIGIGGT CAUGITICCIC GIGGIGCAGT ACCIGACAIG AGCCAGCCAC

MICCCCGAGG GAAAGAATGG CITTIGGTGGC TITIGTTCACA CAGCTGATGC

600 540 480 420

660

6

35

GARAGOGOGA CTCAGARGAA AGATCCTTGA CATTGCCMAA CATGCTGGGC TTGTCCAACA CAGTGATGCG GCTCATCGAG AARCGGGCTT TCCMAGGACA AGTACTTTAT GATAGGTGGG

180

120

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360 300 240 CCTCATTTTA GATOGGCCNC AATATTTAAG ATGGACTGRG GMCCCCARAG ACTGACCCTT

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SEQUENCE DESCRIPTION: SEQ ID NO: 207:

AANCCACTOC ANTITAAACC CCCTCCCCTC CAAGAAAGTT CACAACCGGC CATGGATGAC

	ì			
			GIGCITITICC TCGGIGGGAC AGTIGCIGGC CCTCTINANT TIGGIGIATG TGCTTCCAAG	540
	UCIUCLAGAA ACCATOTICA AGGIAATTAA AAGGICCOTO GOOCGAGCCA GCCTGAGCTT	120	TAICTAAACC TCCAGTCTGA TCTGTATATG CTATCCTAAC TGTTAATTGT AFTAITGAIT	909
v	GCTCACCTTC AAAGTCTATG CAGCACCAAA AAAGGACTCA CCTCCCAAAA ATTCCGTGAA	180 5	ATOTICATIA TCITCCITICA AGGITCATAC TITICAATIT GAIAGAARA AAGITITITI	099
1	GOTTGATGAG CTTTCACTCT ACTCAGTTCC TGAGGGTCAA TCGAAGTATG TGGAGGAGGC	240	U	661
	AAGGAGCCAG CITGAAGAAA GCAICTCACA GCICCGACAC TAITGCGAGC CATACACAAC	300		į
10	CTGGTGTCAG GAAACGTACT CCCAAACTAA GCCCAAGATG CAAAGTTTGG TTCAATGGGG	360		
	GTTAGACAGC TATGACTATC TOCAAAATGC ACCTOCTGGA TTTTTTTOCGA GACTTGGTGT	420	(2) INFORMATION FOR SEQ ID NO: 211:	
5	TATTGGTTTT GCTGGCCTTA TTGGACTCCT TTTGGCTAGA GGTTCAAAAA TAAAGAAGCT	480	(1) SEQUENCE CHARACTERISTICS: (A) INDICH 502 hase pairs	
2	AGTOTATCG CCTOGITTCA TOGGATTAGC TOCCTCCCTC TATTATCCAC AACAAGCCAT	240		
	COTOTITICC CAGGICAGIG GGGAGATIT ATATGACIGG GOTITIACCAG GATATAINGT	009		
70	CATAGAAGAT TTGTGGAAGG AGAACTTTCA AAAGCCAGGA AATGTGAAGA ATTCACCTGG	660 20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 111:	
	AACTAAGTAG AAAACTYCAT GYTCTGCCAT CTTAATCAGT TAFRGGTAAA CATTGGAAAC	720	GARACTGACA TIGITAAACA CACTAAAACA GAAGTACTTA CCTCTTGAAG ATTTAATATA	09
2,5	TCCATAGAAT AAATCAGTAT TTCTACAGAA AAATGGCATA GAAGTCAGTA TTGAATGTAT	780	TAATGGTTGA CATGATACAT GTACATGAAT GGAATGACCA GATGCTTATG GTCTACATTT	120
3	TANATIOGCT ITCITCITCA GOAAAAACTA GACCAGACCI CIGITAICIT CIGIGAAATC	840	TCCTTTATCC TGTTAGTATT ACCTTCCTTA ATCTTTGTTC CTTAACATGC TAAATTCCTC	180
	ATOCTACAAG CAAACTAACC TGGAATCCCT TCACCTAGAG ATAATGTACA AGCCTTAGAA	006	TICAGIGITI ATTITICIAGI GACAGANIGO TAACAITITOT TACACCOTGG CAGAAGGGAG	240
30	CTCCTCATTC TCATGTTGCT ATTTATGTAC CT	932 30	agaaaictigt titiogogiog gtaactaaat tittgagiga aatatcataa gaigagaatg	300
			GAAAGAGGGA GACACAAAGA GITATAACAA AAAAACAATG GITITITIAG CCATITIGACT	360
35	14	\$E	GOCTOTITAA ATAOTOTACA AGACATICAC GITNAACATO ACTITIAGIG AAATAAAATG	420
3	(4) INFORMATION FOR SEQ ID NO: 210:		TOCCATACTA CTATGTOCTT CAAAAGGCA AATGTGCTTT ACTGCCCTAA GGCTAAATTT	480
			TOSTCATTIG ACATCAGAGA TOTTGTAAGT ATTGCACTTA ATACCCACCT ATTTCTCAAT	540
40	(B) YYPE: NUCLELC ACID (C) STRANDEDNESS: double (D) TOPOLOGY: linear	. 40	AGTONTATT TITTGGCTAG CATTINCTTT ACCACTAACC TIGTIGGATA GC	592
	(x1) SEQUENCE DESCRIPTION: SEQ 1D NO: 210:			
45	GTCATTCTTT AMERADAGC TITCCTGTTT AMACCITITC AMAGENGEAG ACCACCITGA	60 45	(2) INFORMATION FOR SEQ ID NO: 212:	
	AGAITCCCCC TAGGGITGAT ATGTGTCTAA TTCATTTTAT AAAAITAIT CITGTCTTCA	120	(i) SEQUENCE CHARACTERISTICS: (A) LEACTH: 938 base bairs	
20	TITITAAAGCT TIGGCIATAT AGTCAGAAAT GICCTAAATA ACAAACTAIT TIGTAITTAA	180 50		
;	TITIAGGGAAG ACTAAAGGGA AGAAAATGA AAACTCAGTC TITIATGTAAG CTCCAAGGAT	240	TOPOLOGY: linear	
	ATTAGGGCTT AAAGGGCTTT TCTAGTTTTA TGAGAAITTG TACTACTGAT TTTTATATAT	300	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 212:	
55	TOCTOTITIT. GAGATGAACA GAICTCTGGG GAAATTGTTG AGTTACAATG GCAITTCACT	360 55	TOGASTICACT TITCCAGCTICA ATGARTCCTA TGTCTCGCGT GCAGGTGGTT GCTTTTCAAT	09
	STGATCCCTC TCAAGCTCAG ATCAGTTCTA TAACCCAATG ACAACCTGTC TCTTTGGTTT	420	OFFICE SCHAME ATTITUTE TATTGGCTCT TGGGAGTTIN CTATGTTIGC TCCTGTGTT	120
09	. ACTOTICCTOT GAMINSTICAG CTCAAGTITIC CCAGAAGTICG TOTOTITATIG ATGAOTICAGA	480	GCCCACCITT AATAAAACA GGCCCAAACA AAAACCATAG CATTCTGAAA CAATAGGGG	180

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CCCACATTOG ACCCAGTATO TCACTTTAAT GGACTTCAAG AAAAAATCTG AATOGGAAA

240

25 40  $\frac{3}{5}$ SS 8 8 GCAGGAGGAA GGGCATGAGC CGAGCTTGAG GAATCCGTGY TCCAAACTCT ACACTCAAGG CGAGACACCC GAGGAGTCCG TTCCTCCCTG GTTACGTGGA CTGTGGAGCT GGTCTCTTGT AGCCTGCCGG GAGAGTGGTG GCATCTRARA GOCTGGTCGT GGACTGTGGT TGGGGGAGGT TOCCCAAGAT GCCCACACAG TCGGAGGTOG ATAACGTOTT TGACACAGGC TTGCGGGACG TOGAACGOGG GOGOTOCTOC AGOTTOCGAG TOCAGOCAGO CTGGGOGGGG GGCGCGCCCC BOURDETT TIRACCOTOT GEOCECETETE ETGTGECKGE GTGGGEATEC CECGGGGEAG AGAAGAAGTC TCCCTOGCCA TTCTCAGACG AGTGCATCCC ATGGGAAGTG TGGACGGTCA RIOCMCIGCG CAACICIGGI GGCGAIGGGC 10000CAGAI GICCTIGGAG ITCIACCAGA GOCTICAGOOC COTOCOGAGO TTOAAGCOTA CCTOCOGAGO TCOCACCAGO GOCOTGAGGA FIGAGAAACT CIGCGAGAAG ATCATCAACA TCOTGGAGGT GATGAATCGG CATGAGTACT AGGIOCAIGI GGIAGCCCIG GCCACGGAGC AGGAGCGGCA GAICIGCCGG GAGAAGGIGG E SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 213: (B) TYPE: nucleic acid (D) TOPOLOGY: linear (C) STRANDEDNESS: double (A) LENGTH: 1079 base pairs 660 600 540 480 420 360 300 120 S

20 25 5 5 S GGTGTCCTTT CCCTGTCAAA CTCATTAAAA ATTCCTTT GCTGAAACCA AACCAGGCTT TTAAAAAACCT GTGTAGAAGA AAACCAAAAA ATCCTGTGTG CATTITICATT GAAGAGICIG AIGACTIGCT AGCGITTIAT CATTOTGACC ATOGTGATGC CTCATTTGCA TGATATGTAC CTTGTGTTTA ATGTGAAATA TECHGGAACE TRATTITIGET AMAIGNETGT TETICITIGIGA TETIGITIGIAE CTEACAGCAC TTATGACAGG CACACAGAAA CCATAGCATG GYCTOGCTTT CAGAAAAATGC CTCTCATCTT AAGTACATGC CTCCACATAA TGCGGTGCTG TCCATCTCGG CAAATACTGG CCAAGTCCCT TIGITICITG TOGCGAMATG TOGCTTTCAA ATTAMAMIGM CCTTTTCTTC TTKGMAACTT ACATTTAAGR ATGTATTAGT TACAGAAATT ATATGTCTGT GTATGTGTCT CTACTCAATI AACCAGGAGA TACAAAGAAG TCTCAGTAGT AATCTTGTTC ATGTGCTTTT TITICITITKGA CIKGIATAAI TAAGGGITIG GAAAGATICA TAATIMIGAG AGAGGITIGG TCACACTAGG AATGTATACT CCACACATTT TATGCCATAT AATGGTGTGT TITCTTAATT TITTICIGIA AGCICAAIGI ACAGCCAGCT 900 938 840 780 720 660 600 540 480 420 360 300

(2) INFORMATION FOR SEQ ID NO: 213:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 214:

(C) STRANDEDNESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 3791 base pairs 20

(2) INFORMATION FOR SEQ ID NO: 214:

(i) SEQUENCE CHARACTERISTICS:

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AGCCACCACT GOGATOGOGA ATAAAGTTGA GAACATGAGT TTGGGCTGAA AAAAAAAAA OCCIOTACIO ICICIOCATA GOOCIOGIOG GUICOCCOTI CITICIOCAC ICIACAGAAG

1079 1020 5

TIGGTITIGI GGIIGCCAGC CICAGGICAI CCITITAAIC ITIGGIGACG GIICAGICCI

960 900 840 780 720

GCTCTGGAAC CTGCTCTGGG TCATTGGTGA GACTTGGAAG GGGCAGCCCC CGCTGGCTTC GGGAGCTCCT TGATGGCTCC CAGACCTTGG CTTTTGGGAA TIGCACTTTT GGGCCTTTGG CCACCACCAT GCGCAGGCTC ATCAAAGACA CCCTTGCCCT CTGAGCGTCG CTGGATCTCT TOCAGECETA CETOTACAAG ATETECETTEE AGATEACTGA TGECETOGGE ACCTEAGTEA

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GYYYYYYYYYY GCCGGCGGG GCGGCGLLGG GGCCCLLGGG GCCYWGYGCG CYLLCYGGGCC GOGGOCCAGA CCAACTOCAA CGOGGCAGGC AAACAGCTGC GCAAGGAGTC CCAGAAAAGAC

CGACCAGTTA TITGACGACG AATCGGACCC CTTCGAGGTG CTGAAGGCAG CAGAGAACAA AGCCACCATC ATOCCTOGGC ACTTACAGGA AGGCTTCGGC TGCGTGGTCA CCAACCGATT TRANSCASSIC SCIPTINGSOT COSCOCOGOC COCTOCAMIC COTOGAGGAA COCOCOCOCO

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CCCGTGGCGC TTTAAGAAAG AAGGAATAAG ACGAGTTGGA AGAAGACCTG ATCAACAACT COCAAGAACC COCTOCCCC CAGCOTTOGC GTOOTTGACA AGAAAGAGGA GACGCAGCCG

> 360 300 240 180 120

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TCAGGGTGAA GOGAAAATAA TTGATAGAAG ACCAGAAAGG CGACCACCTC GTGAACGAAG 420

TATTGACCGA CCTATTCGAG GTCGTGGTGG TCTTGGGAGGA GGTCGAGGGG GCCGTGGACG ATTICGAMAAG CCACTTGAAG AMAAGGOTGA AGGAGGCGAA TITTICAGTTG ATAGACCGAT 540 600 480

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TOGRATICOCO CORGORGATO GATTIGATIC TOGTOGORAR COTGRATITO ATRIGOCATAG 660

TOGRANGIGAT AGAICTICIT TITICACATIA CAGIOGCCIG AAGCACGAGG ACAAACGIGG AGGTAGCOGA TCTCACAACT GOOGAACTOT CAAAGACGAA TTAACTGACT TÖGATCAATC 780 720

840

AAATOTOACT GAGGAAACAC CTGAAGGTGA AGAACATCAT CCAGTOOCAG ACACTGAAAA

OTOGRAGOCT ATTICARANTA AGGRECOGGE ARANGTAGRA TITRATRITCE GRARACCIAR

TAAGGAGAAT GAAGTTGAAG AGGTAAAAGA GGAGGGTCCA AAAGAGATGA CTTTGGATGA

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TGAAGSTGCT GATGSGCAGT GGAAGAAGGG ATTTGTTCTT CATAAATCAA AGAGTGAAGA	1020	ATTIAGCIGA TIGGITCICA CATATACTIC TAAAAGAAAC TITTATGITA TAAGAGTIAC
GOCTCATOCT GAGAITICGS TIATIGGACCA TCATITICCCG AACCCAGCAA ATGAIATIAAC	1080	TITITICANIA AGAITIANTA ATCICAGITA CCIACIANTIC TGACATITIA GGAAGGAGGI
OTOTCHACTIC CHANTCHAIT THEGRANCT TECCOCCCA GRACTICOCG CCAGGGAGG	1140 5	AATTOTTTT AATGATGGAT AAACTTGTGC TGGTGTTTTG GATCTTATGA TGCTGAGCAT
ACCARGOTICGA COTGGGGGTG GTGGGGGGCCC AACCGTTGCC AGCAGGACCG ACAAGTCAAG	1200	GTTCTGCACT GGTGCTANTG TCTNATATA TTTTATATTT ACACACATAC GTGCTACCCA
TECTTCTECT CCTGATGTGG ATGACCCAGA GCCATTCCCA GCTCTGGCTT AACTGGATGC	1260	GAGATTAAIT TAGTCCATAT GAACTATTGA CCCATTGTTC ATTGAGACAG CAACATAÓGC
CATAAGACAA COCTOOTTOC TITTOTGAACC CITCTOTTCA AAGCTITTGC ATGCTTAAGG	1320	ACTICCIMANT CAGIGIGITIT AGACTITICA AGIANCIMAC TCATTICCAA ACATGIACCA
MITCCAAAGG ACTAAGAAT TAAAAAAAA AAGACTGTCA TTCATACCAT TCACACCTAA	1380	TGITITIATAA ACCICITGAI ITCCAGCAAC ATACTATAGA AAACACCTGC TACTCAAAAC
AGACTGAATT TTATCTGTTT TAAAAATGAA CTTCTCCCCC TACACAGAAG TAACAAATAT	1440	ACAACTICIC AGIGICAICC ATIGCIGICG TGAGAGACAA CAIAGCAAIA ICIGGIAIGI
COTACTCAGT TITICTATITA GAAATGTAIT COTACCAGGG ATGITITICAT AATTITICAGA	1500	TECANGCITT CAAGATAGCC TGAACTTAAA AAGTTGGTGC AITHGTTGTA TCTGATGGAT
CATTAIGCAT TCTTCATGAA TACTTTIGTA TIGCTGCTIG CAAATAIGCA TITCCAAACT	1560 20	ATAAATTIIGC CICCIAGIIC ACTIIGIGIC AAGAGCIAAA ACIGIGIAACC TAACITIICIC
TGAAAITATAG GTOTGAACAG TOTOTACCAG TITAAAGCTT TCACTTCATT TOTOTITIT	1620	TIAITIGGIGG GIAATAACIG AAAATAAAGA TITIATITICA TGCTCACTIC TIAAAAGICA.
AAITAAGGAI TTAGAAGTTC CCCCAAITAC AAACTGGTTT TAAATATTGG ACATACTGGT	1680	TANANACANT CANATAGGRT CANGITTAIT GICATGTGIT TCCTGGKTTC TGACCTGTGT
ITTAATACCT GCTTTGCATA TTCACACATG GTCAACTGGG ACATGTTAAA CTTTGATTTG	1740 25	GCACACCCCT GTGTGTTAT ANTITTTAAA TTGAATTTTA TATGGGGTTT TTATTTGCTA
ICAAAITITA IGCIGIGIGG AARACIAACI ARAIGIAITI TAACITAGII TIAARAITIT	1800	AAAACCAGGC TGITGAATCA CATTTGGGAA GGGTACTTAT CTTAATGACT AATGACTTAA,
CATTITIGGG GAAAATCIT TITICACTIC TCATGATAGC TGTTATATAT ATAIGCIAAA	1860 30	TIGGGAAAGT TGAAITCTIG TAAAATACAA AATCCAAGGA CITCTIGGGA TITLAATCTAA
TCTTTATATA CAGAAATATC AGTACTTGAA CAAATTCAAA GCACATTTGG TTTATTAACC	1920	TYGICACTIC NITAGGCAGA THCACTITIT TGGATAATGG AAAGTTAAGC ATACCGAATG
CTIGCICCIT GCAIGGCTCA TIAGGITCAA AITAIAACIG AITIACAITI TCACCIAIRI	1980	CTACTITITIGG TIGACAAAGG GGCCTAAING TCCGGGGGGA AATCCCTAAC NGGTAAGGNT
THACTITITH AMIGCITICAG ITTICCCATIT TAMAMICINA ACTAGACATC TIMALTICOTG	2040 35	CCCAACTATG G
AAAGTIGITT AAACTACTTA TIGTIGGIAG GCACATCGIG TCAAGTGAAG TAGTITTAIA	2100	
GGIANGGGIT TITTCTCCCC CITCACCAGG GTGGGTGGAA TAAGTTGATT TGGCCAATGT	2160 40	(2) INFORMATION FOR SEQ ID NO: 215:
GTAATATITA AACTGTTCTG TAAAATAAGT GTCTGGCCAT TTGGTATGAF TTCTGTGTGT	2220	(1) SEQUENCE CHARACTERISTICS:
GAAAGGTCCC AAAATCAAAA TGGTACATCC ATAATCAGCC ACCATTTAAC CCTTCCTTGT	2280	(A) LENGTH: 1334 base pairs (B) TYPE: nucleic acid
TCTAAACAA AAACCAAAGG GGGCTGGTTG GTAGGGTGAG GTGGGGGAGT ATTTTAATTT	2340 45	(C) STRANDEDNESS: double (D) TOPOLOGY: linear
TIGGAATITIG GGAAGCAGAC ACTITIACIT ISTAAGGITIG GAACAGCAGC ACTATACAIG	2400	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 215:
AAATATAAAC CAAAAACCTT TACTGTTTCT AAATTTCCTA GATTGCTATT ATTTGGTTGT	2460 50	CASTGCTCGC TOCTGCTCGG GGGGCTGCGG CCCCGGGGCGT CGCCATGACC AGTGAGCTGG
AAGTIGAGTA TICCACAGAA AGTGGTAATT ATCTCTTCTC TCTTCCTCCA TTAGAAAATT	2520	ACAICTICGT GGGAACAGA CCCTTAICGA CGAGGACGTG TATCGCCTCT GCCTCGATGG
AGGTANATAA TGGATTCCTA TAATGGGAGC ATCACCACTT ATTAAAACAC ACATAGAATG	2580	THACTOGGTG ACCOACCGGG TGGCCCTGCG GGTGCGCTCG GGAATCCTGG AGCAGACTGG
ATCAATTAAA AAAGTITICT AGGATIGICT ITTAITICIGC CACATTAATT GALAAACAGF	2640 55	CGCCACGGCA GCGGTGCTNC AGAGCGACAC CATGGACCAT TACCGGACCT TCCACATGCT
GAAGGAATTI TTAAAAATT TTTAAGAATT GTTTGTCACG TCATTTTTAG AAATGTTCTA	2700	CGAGCGGCTG CTGCATGCGC CGCCCAAGCT ACTGCACCAG YTCATCTTCC AGATTCCGCC
CCIOTATATO GTANGICCA GITTIAAAA TATIGGACAT CITCAATCIT AAACAITICI	2760 60	CICCOGGAG GCACTACTCA TOGAGAGTIA CTATGCCTTT PATGAGGCCT TTGTTCGGGA

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GGTGCTGGGC AAGAAGCTGT CCAAAGGCAC CAAGAAAGAC CTGGATGACA TCAGCACCAA

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GOGGRACITY TRACARGEET CRECIBECTE RERECCERGR CECRCITECT GETECRACIAC

660

600 540 480

720

CAGCCCCTCC AGCTGACCAC CCTGTTGGAC CAGTACATCA GAGAGCAACG CGAGAAGGAT

S

STACSGATCT TAAGGACTST GCCATTCCTG CCGCTGCTAG GTGGCTGCAT TGATGACACC

CCCCTTTTTG TGGGGCACAA CATGCTGCTG GTCAGTGAAG AGCCCAAGGT CAAGGAGATG RICTICCCCA ACCAGGICCT GAAGCCCTTC CTOGAGGATT CCAAGTACCA AAATCTGCTG

ANCONCAGOA GGCAGGGCTT TANCAACTAC TOCAAGCTOC COAGCOTGCC CONGGTGCAG

45 6 35 ઝ 25 20 55 8 5 5 S CTUCCATCTC CTCCCAGCCC CCCACAGGAG GAGATAGGCC TCATCAGGCT TCTCCGCCGG 2) GIGINCINGIG GCICCTIGGG AACIGAGACA TAICICAGGG AATGGIGICI GIGCICAGCC GICTOTCCAC ICCCICGAIG GCITCCGACA CCAGGCCICI GOGACCGCIA CAIGGGCACC CTOCCOCTCC GACCACTOGC CACTCAGCGA COTOCGGTTC TTCCTGAATC AGTATTCAGC GACCCACAAT AAAGATGTCA GAGACCTGTT TGTGGACCTC GTGGAGAAGT TTGTGGAACC TOTOGRACOTO CACARGRAGO TOGTOTOCAO TOCTOTOCO GRARAGOTOG GOSTOTTOTO AGACAAGGAA TITICTCCAGG ACTIGAAGGA GCTCAAGGIG CTAGTGGCIG ACAAGGACCT GATCCAAAAC TOGACCCTTG GAGCCGTCGA CTCACAGATG GATGACATGG ACATGGACTT AGGGAAGAAA AAACTGCAGT ATCTGAGCTT CGGTGACTTT GCCTTCTGCG CTGAGCTCAT GETGETAGAG GAMATGEGGG GETECETTGET GGACAATATT CAGCAACACT TECTECTETE AACAGGCATC ACCCTCAAGA GCTGCCGGAG ACAGTTTGAC AACTTTAAAC GGGTCTTCA GIGGCGGGGA IGCIGCGAGG GGGICICCIG CCCCAGGCGG GCCGGCIGCC TACCCICCAG GCTGACAATA AAGTTGCTCT GAGTTTGGAG ACTGGTCCTC GCTCCGGGGA GCAAGTGGG TGAGATGGAA GCCAACTTCA AGAACCTGTC CCGGGGGCTG GTGAACGTGG CCGCCAAGCT TGACCGGTTG GCCAGGGACT ATGCAGCCAT CGTCTTCTTT GCTAACAACC GCTTTGAGAC CAGAAGCTGA TOGCTGTGAC TGAATATATIC CCCCCGAAAC CAGCCATCCA CCCATCATGC AGTGCAGAGO ACAAGCTTCT TATGCGACAC CAGCTGCGGA AACACAAGAT CCTGATGAAG GAGATAGCAG CAGTITITCCA GGACAACCGA ATGATAGCCG TCTGCCAGAA TGTGGCTCTG ACTOTOGOT ATOGOTOGAA GGOTOTTACO COCCACOGTO OTOTGATOGA CTTTCAGOGG INFORMATION FOR SEQ ID NO: 216:  $\Xi$ (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 216: SEQUENCE CHARACTERISTICS: TOTOCCTOTO TCTGTCTCTO AGCACCTGGT GTCCGTGTAC AAGGATGGAT GCCTCCTGCG CCTGTATCAT GACTGAGGTG CCTCCCAACG CTCCGCCCAC (B) TYPE: nucleic acid
(C) STRANDEDNESS: doubl
(D) TOPOLOGY: linear (A) LENGTH: 1511 base pairs STRANDEDNESS: double 1320 1140 1080 1334 1260 1200 1020 960 900 360 8 240 180 660 600

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8 2 6 330 25 20 8 55 7 GCCTCTTGCT GCACGGGCTC CTGAGCCACC CCCTTGGGGC ACAACCTGCC ACTGCCACAG TOTTTAGCCA GCCCTGCGCA TAAATACACT CTGCGTTATT GGCTGTGCTC TCCTCAATGG TITIOTCAGG TIGICCTIOT TIGGATCCCT CAACTAGGIG ATAAGCACIG GAGGGGGAIG GACAAAGITC ACGTAGCAGG TCTAGGCAAA GACTGGGCAA TTGAGCAGAG GAGACGGACC AGGCCTTACT TITICCICCCA CAAAGGAGIC GCAGCCACGC TAGCICIGAC (2) INFORMATION FOR SEQ ID NO: 217: aaaatgaaat a TOTTAGTICT CCACCCICGA GOTOTACGCT GIGAAAAGTT TGGGAGCACT GCTTTATAAT AGAGOTTOOT CATCAGOTTA COCAGAAGTO GOTOCCATOC ACCATOCAGG TOTOCTTOGA CAGAACAGGG ACTTGAAACCA AGCCCTCTGC TCTGAAGACC GCGTCCTGAA TTTCTTCACT TCACAGAGAA GGAGCAGAGA AATTAAGTOG CTTOCTCAAG GTCATGCAGT TAGTAAGTOG CTITIVICCIU AGIUCIVIAC AIGIATIAGI ICCITTACIO CIUACCACAI IVIACCCAIT TROCTCAACC AAGCAGTTOT GCTGAGAATG OCACCTOGTG AGAGCCTOCT GTGTGCCAGG CAGGITCCAAG GCAGAGATCC TGAAAAAGATA GGGCTATTGT CCCCIGCCIC CTIGGICACT COCCTICOTO CITAGITITIC CCAACITOGG ACGIGATAGG AGCAAAGICI CICCATICIC ATATTOTCAG GTATAGOGCC ACTTOGAGAT GCAGAGGATT CCATTTCAGA TOTCAGTCAC GACATOTOGA AGAACTITOGO GICOGOGAGI GIGITITOTCA CIIOGITITIC ACTAGIAAIG TOTOTOATOT COGCOMATOG GAAGCOAGAT COTGACACTG TTCCCGGACTC GTAGCOAGCC (1) SEQUENCE CHARACTERISTICS: ACCRYGAGSC GGRCCCCTTC ACCTTGGCTG GGCTGGTCCT GGTCCTTAGG SEQUENCE DESCRIPTION: SEQ ID NO: 217: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENOTH: 642 base pairs TIGCCACIGI 1500 1440 1380 1320 1260 1200 1140 1511 1080 1020 960 180 900 240 120 60

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			CIOCAGITAC CATARITITIC CAUGITIGIG GAAITGATAT TGAAATAGCA GGGCTAAGGÀ
	ACCOGCUTIG GACGIGITIC ITTAACCICA TCCATATAAT AGGGCCGTGG GAIGGITIGTA	300	ATTACTOCICA AGITTTAGCC TOTGGGTAAT ACCTTAGGGT TATTTAAATA TITGTAATTT
<b>v</b>	GAGGTAAAGC AGGATGATGG TOTTTTAAGA CCAGAGCTTG GGACCAGGGC TCCTACACCT	360	TATTTAAATG TTCATGAATG TTTGAAAGGA ACAAAATTAT CAGGGATGGC TCTTTGGCAT
)	AATTITICICT CCTGGTAGCT GAACAAAGGT CTAAATTAGC TTAACAAAAG AACAGGCTGC	420	GOSTICTIALT TICACCTICT TITICIONAG AAAAAAGAAC AATGICITIAA TGIAITITIA
	COTCAGCCAG AGTICTGAAG GCCATGCTTT CAGTTTCCCT TOTTGACAAT TGCTCTCCAG	480	AAGITITING TANAGITICH AATHOCAATT TIAANAAAG T
01	TTCCTATGAA AGCACAGAGC CTTAGGGGGC CTGGCCACAG AACACAACCA TCTTAGGCCT	540 10	
	GACCTOTGAA CACCAGGGGG TITGTOTOTOT OTTETTTTC TETECTTGCC GAACTITICTC	009	
ž	AATAAACCCT ATTTCTTATT TTATATTTAC GINGGROCTG GG	642	(2) INFORMATION FOR SEQ ID NO: 219:
2		•	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1080 base pairs (B) TYPE: nucleic acid
70	(2) INFORMATION FOR SEQ ID NO: 218:	20	
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1241 base pairs		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 219:
ž	(B) TYPE: nucleic acid (C) STRANDEDESS: double	25	TOTITATIONS ACCIMADAGA TACACACATG CACACACACA TACATATICA TTCATTCATT
3	(D) TOPOLOGY: Linear	}	CATICAAGIG GIGITICCAG TGICIGIGIG TCACTGITIA TGCAGITICC AITITCCCAGI
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 218:		GAATTATGAG TGGAGGGAA CTTTTCTAAC CAGATTGTCT TTTCAGAACA AAGACCKGGG
30	GORCCCACTG TICCATITIA IGCTAATAGA TICCATICTA GGGCCCAGCC GICTCTIGAC	90 30	RATTGAGGAA GAGTTTTGGAA AGAGGGAGAG GCAAGGAAAG AGAGCTTTAA ATTGAAAGGT
	TGATGOTGTT CCCTTTAACC CTTGGCATGT ATAATAGAAT TTTGGTGAAT GAAAGAACC	120	TAATTICCTA AGAGGAACCT GGGCTGAATG ACTACAGTGT TATACCCTCC AATCTTTGCA
	AAAIAGGCCA GAIAGTCCCC CCAGGCCCTG ATATCCATAA AAGGCTTGGG AATGCATTAT	180	GOTGGGCATG GAACACTGCT TGTATCACTC TGTGCACGGT ATAAATCCAT ATATCCACAA
35	GIAATIGICC TIAGICITIT IGIIGITITA GAAAAAAA ACAAGAIGGG CICAGAIGGA	240 35	AAACACAA CCATCCATCA ACATATACAT GGTTTTGGGAT GAGCAGGTCA ATAGTTTTGA
	TGCCTACGTA AAAATGGTTC CTAGCTGTGT ACTCATAACT TTTCTTTGAA TTGAGTAGTG	300	GAGGGAGTIT GITCCTITIT TITTICICATI APACTCTIAA ATTGTTGTCA GTTATCAAAC
. 6	AAAGGAAGGA GGAGGAAAGG AAATTAAATG TOCTTOTAGT ATTOTOTGGA CTCAAGTOTG	360 40	AAACAAACAG AAAAATTGTT TGGGAAAAAC CTTGCATACG CCTTTTCTAT CAAGTGCTTT
!	ACATATERGA TAATAACCTA TATTGAAATG CCAAGAATTG TATCTGAAAC AAGRGAACAG	420	AAAATATAGA CTAAATACAC ACATCCTGCC AGTTTTTTCT TACAGTGACA GTATCCTTAC
	TITICACACAT TTATCATGCC TTCATATTAC ATATTAACTG AAACCAATTA ATAAACATAT	480	CIGCOATITA ATAITRAGCCT CGTATTITIC TCACGTATAT TTACCTGTGA CTTGTATTTG
45	GAANTATICA TTGCACAAGG CAAAGGCACC TAAACCTETT GTTTCTTTTT CTACATAGCA	540 45	TTAITTIAAAC AGGAAAAAA ACATTCAAAA AAAGAAAAAT TAACTGTAGC GCTTCATTAT
	GAAATTGAIT TITTITIAT TITTITAGGG GAACCTATAT AATTATGACC CAGTGATGTC	009	ACTATTATAT TATTATT ATTGTGACAT TTTGGAATAC TGTGAAGTTT TATCTCTTGC
20	ITTIOGICAC TIAAGCITAI GAATICAGGI TACAATICAG TICAITCIAG AIGGITACIA	099	ATMINCITITA INCOGNAGITA TINCOCCTIA AMAMINCGAN ANIMANITITI ACANGGITITC
	CCTTGAAAAG GATGTTGGTG CCTTATGTGA CAGGAGCCAG AGCCTGCTGG GAATAAACAA	720	TETITIGICI GGAGGGIAA TIGAICITICC TAAGAATGAT GITTIGITITT TIGGGGTTTT
	AGCAGATTCA TGCCAACACC AACTGGTAGC TTTAGTGGCA GATGGGAGTG GTCACAGACT	780	TGTTGTTTT TTTTTAAATG TTACCAGCAC TTTTTTTGTA AGTTTCACTT TCCGAGGTAT
22	CCCAAAATOT GGGGCTTTGG ATTTCCACAC CATCCCACGT GTGTGACATC TTCCTCTTTC	840 . 55	TGTACAAGIT CACACTGITT GTGAAGITTG AATATGAAGG AATAATTAAA AAAAAAAAA
	ACACTICITICA TIGATARITIG AMANTENGA ANTICICITI GANTITIGICIT ATACCATICAG	006	AAACCHOGGG GGGGGCCCGG TCCCAFTTGGN CCCAAGGGGG CGGTTACGGG GTCACGGCCG
8	CACATICTTA TGACAACATA ACAAATAGTT CATAATGTGA ATATTAGAAA CTGTTACAGC	09 096	

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SEQUENCE CHARACTERISTICS

120 180

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INFORMATION FOR SEQ ID NO:

50 45 8 ઝ 30 25 20 15 5 TCTGAGCTTA TITTTCCATT TGATATICAT TGATATCATG ACTTCCAATT GAGAGGAAAA TITTICICAGA AGGATKAGAA ATCAGCACCT GCGTITTIAGA GATCATAATT CTCACCTACT GIGCCAGITA GIGCCITCGG IGIAAGAICT ICICATCAGC CCICAAITTG IGAICCGGAA AACCCAKGTG CATTICIGCA TCTCCTGGAT TAGCCTTTSA CATGITGCTG RCTCACATTA AATTOTOTOC TUACTACTOT CCAGTAAGGA GGCCCATTGT CACTTAGAAA AGACACCTGG GAGGICTITT AAGTCTCTTC CCTTTGGTTG CCCACCTGAC MATTTTATTA AGTACATTTG AAGTECTECA GETGGTECTG CIGCTAGTAG TGTTTGGYTT ATTTTCCATE CCAGTICTGG AATTCCACAG AAGATATCAT GTCTTTGTCC ACCCAAGGGG ACTACATCAA TCTACAAACC GTCAAGGAGA ACAAAAATAT TAATGAGGCT ATGAGAGTCC TCATTGAAAA GATGATGAGA COGGASCAGA TIGACCOGIT CAGTAAAGAG AACGGTITCA CAGGTITOGAC AGAAACAICA GAGCCGGTGC CCTGCCTCCTT CTTGGCCAAC AAGTGTGATC TGTCCCCTTG GGCAGTGAGC TTCAGCAACA GCCAGAGGTG GAAACAGGAC CTAGACAGCA AGCTCACACT ACCCAATGGA TIGINITATE GGGAIGCCIC TOCCIGIGIT AFTAIGITIG ACGITACCAA TOCCACIACO ATAGTOCOGC TICAGCTOTO GGATATIOCA GGOCAGGAGC GCTICACCIC TAIGACACGA AMIGIACATA TATITICAGI GGATTITIGCI CIGAAGGITIC ICCAGIGGIC IGACIACGAG TECCTATITE ATTICACITY ATCCCTATTA ATTITITACA CICAAATITY ATTAAAGIAT TGAATTGAGG GCTTAAAGAT AAACATATGG GRTTGGAGTT GTOTGTCCAT AGGGTTTCAC IGGCTACTAG AAGTOTCCCA GAAGTCACTG TATTTTTGAA ACTTCTAACG TCATAATTAA IGAGATOAAA TOTOATTIOC CAAATTIOTT OTAGGOOGIT GITTOAGATT CITICIGIOT Ĕ ACATCTGATT CTGGAATGCA GAAGGAGGG TCTGGGCATC TGTGGATTTT SEQUENCE DESCRIPTION: SEQ ID NO: 220: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear LENGTH: 1258 base pairs

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SEQUENCE DESCRIPTION: SEQ ID NO: 221:

(A) LENGTH: 1693 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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8 55 50 45 6 35 30 25 20 5 0 ATAGAAGAGG CCATATATAT TOCCICCTTA TCCTTGAGAT TTCACTACCT TTATGTTAAA TATTCCCTTT AAGAATOTTT ATGTATGAGT GTGAAGATGC TAGCGAACCT ATGCTCAGAT GAAAGITIGA CCAAATITGI TITITIGITG TIGITGITGI TITGAATTIG AAATCATICI ACTOCCCTAC TCAAGACCCC CTAAGACTGG TAGAAATTAA AAGGATTTCA AAAACTTTCT TTATOCAATO GTTTOTTOAG ATACOGACIT GATGOTOCTO TTTAATCAGI TTGCTTCCAA TATIGAGGAA AGGTATICIT CYATACAACT IGITTIJAACC TITGAGAACA TIGACAGAAA GAGAATICAG GATAGITTIG TITAAAITICT TGCAGATIAC AIGITTITIAC AGIGGCCIGC TTATTTAATT TITTÄGGTAA TGCCTATCTC TTGGTCTATT AAGGAAAGAA GCAATCAGTA TICAANIACC CAAAIIGIGA TAGCAIAAAI AAAGIATIIA TITIAIGCCI CAGIATAIIA GCTGGCACCG TIGCTICCIC TITGGGAAGA GGAAAGGGIG TGTGAACATG GCTAACAATC GCATATTGCA GAACAGCTCT GAGAGCAACA GTTTCCCCATT AACTCTTTCT GACCAATAGT GAAAAAAAT CATTTATCC GICTTTTAAG TATATGTTTA AAATAATAAT TTATGTGICT TITIGGITICAC TIGGCITIAAGA GGITTICIICAG AATATCTIATIG GCCACAGCAG CATACCAGTT TOSCATTATO TITTITACAAA ATAATGACAT ATGICACATG TITGCATGIT TOTTITGCTTG AGAGCAACAT TOTOTTATTA AAGCATAGTT TATTTCACTA GAAAAAATTT AATATCAAGG TITICCAGIGT TGAATCTCAC ACACIGTACT TIGAAAATTT GTATTTATTG TGCTGTGTCT GGTCCTAAGT GGAGCCAATT AAACAAGTTT ATTICTITAG TAAGAATGIG TIAAAAITAC AATGATCITT TAAAAAGATG ATTICCTITICT TARACCTACC AGCARACTAG GATTIGTGATA GCAATGAATG GTATGATGAA TCCATCCTAA TAGGAATGAA ATTAATTITG TATCTACTGA TAACAGAATC TGGGTCACAT ATTAMAGGCT GITTGCACCT TIMAGGACCA GCIGGGCIGT AGTGATICCI GGGGCCAGAG AAAACITGAT CCACATCACA CCCIGITITAT TITICCITAAA CAICITGGAA GCCTAAGCIT CACAMINIAI GAAATAGIAC CCICIAAAAA AGAGAAAAA AAAAICAGGC GGICAAACIT ATTCATOGTA AGTOTOCOTT CACOTGITAC AGAGTITICAG ATOGGTOACT GATAGTATGT ITGAATTITT GAACAGCCAG TIGACCAATC ATAGAAAGTA TTACITICIT TCATATOGIT CTGAGAATCA TOTGGCAAGT GTGATGGGCA GTAAAATACC AGAGAAGATG TTTAGTAGCA ACTATTACAT ACTICATTAC TAGGAAGTIC TITITAAAAT GACACITAAA ACAATCACIG CCTTCCATCC TGAATAACGA 1620 1140 1080 1560 1500 1440 1380 1320 1260 1200 1020 960 900 840 780 720 660 600 540 480 420 300 180 120

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GITTICICITG

TCTTGGGCAT CAAGANTAGT TCCAATTTTT TGGGCCGGGG CAGGGTGG

1258 1200 1080 1020 960 90 840 780 720 660 90 540 480 420 360 300 240

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 221:

	WO 98/39448	PCT/US98/04493	WO 98/39448
			450
	acticiciat aaticitaaa aictgigaaa gaataaaag tggaittaal itaaaaaaa	1680	(2) INFORMATION FOR SEQ ID NO: 223:
8	AAAAAAAAA AAA	1693	
٠			(B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear
10	(2) INFORMATION FOR SEQ ID NO: 222:	10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 223:
	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1196 base bairs		TCAGGGAGGT GOCAGGAAAG GCTTGGAACA GCTGCCGGAG TGACGGAGCS GCGGCCCCCC
	(B) TYPE: nucleic acid		
15	(C) STRANDEDNESS: double (D) TOPOLOGY: linear	. 15	COGGINGOSC INGARGINCA ABCITICAGG TARCOGOCOS CAGAGOTINA CCOAGGOTOT
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 222:		GGACATCCTG AGCCCAAGTC CCCCACACTC AGTGCAGTGA TGAGTGCGGA AGTGAAGGTG
			ACAGGGCAGA ACCAGGAGCA ATTICTGCTC CTAGCCAAGT CGGCCAAGGG GGCAGGGCTG
20	ACCCTICACT CACCCACCC CTCCCCACN TGCCTICGTG GGGAAGGAA AAGAITTGT	60 20	OCCACACTCA TICATICAGGT GCTIGANGGC CCTIGGTOTICT ACOTICTITIGG AGAACTIGCTIG
	AAACCCCGGA GCGAGGTTCT GCTTACCCGA GCCCGCTGCT GTGCGGAAGAC CCCCGGGGTGA	120	GACANOCCCA ATGITAGAGA GOTGGOTGAG AGTGACTITIG COTCTACCTT CCCCOTGCTC
•	AGCCACCGTC ATCATGTCTG ACCAGGAGGC ANACCTTCA ACTGAGGACT TGGGGGATAA	180	ACHIGICATION OF AN ACTION OF A CONTRACT AND ACTION OF A CONTRACTOR AND ACTION AND ACTION AND ACTION AND ACTION AND ACTION AND ACTION ACTION AND ACTION AND A
25	GARGGARGOT GARTARATTA AACTCAAAGT CATTGGACAG GATAGCAGTG AGAITCACTT	240 25	ארשותווות מוואותפתאר אוארתיותאה וארוואתיות שאתיההתתא זכווההורה
	CAAAGTGAAA ATGACAACAC ATCTCAAGAA ACTCAAAGAA TCATACTGTC AAAGACAGGG	300	CTANCAGAGG CTCAGAAGAA TANGCTTCGA CACCTCTCAG TIGTCACCCT GGCTGCTAAA
	TOTTCCAATG AATTCACTCA GOTTTCTCTT TGAGGGTCAG AGAATTGCTG ATAATCATAC	360	GIAAAGIGIA ICCCATAIGC AGIGITISCIG GAGGICITISC CCTGOGIAAT GIGGGGCAGC
9	TCCAAAAGAA CTRECTAATIGG AGGAAGA TGTRAKTIGAA GTTTATCAGG AACAAAGGGG	30	TGGAAGACCT TOTGATTGAG GCTGTGTATG CTGACGTGCT TCGTGGCTCC CTGGACCAGC
			GCAACCAGGG GCTCGAGGTT GACTACAGCA TCGGGCGGGA CATCCAGCGG CAGGACCTCA
ţ	GOSTCATICA ACAGITIAGA TAITCITITIT ATTITITITIC TITICCCICA ATCCITITIT		STRUCKATTRC COGRACCETS CAGGIATIGST STOTEGOCTIO TRAGSTOSTS CTOTCAGGCA
દ	ATTITIADA ATACTICTIT IGTAATOIGG IOTICAAAAC GGAATTGAAA ACTGGCACCC	540	TTGAGGAGCA GOTGAGCCOT GOCAACCAAC ACAAGGAGCA GCAGCTGGGC CTGAAGCAGC
	CAICICITIG AAACAICIGG TAAITIGAAI TCIAGIGCIC AITAITICAIT ATIGITIGIT	009	AGATTGAGAG TGAGGTTGCC AACCTTAAAA AAACCATTAA AGTTACGACG GCAGCAGCAG
9	TTCATTOTIC TGATTITIOG TGATCAAGCC TCAGTCCCCT TCATATTACC CTCTCCTTT	660 40	COCCADODAC ATCTCAGGAC COTOACCAAC ACCTGACTGA GCTGAGGGAA CCAGCTGCTG
	TAAAAATTAC GTGTGCACAG AGAGGTCACC TITTTCAGGA CATTGCAFTT TCAGGCTTGT	720	GONCOANCOA GOOCCAAAAAAAAAC CTCAAAAAAAGC AAGGGGGCTCC GAAGGAAGCCC
	GOTGATAAAT AAGATCGACC AATGCAAGTG TTCATAATGA CTTTCCAATT GGCCCTGATG	780	CARCAVITICS INCLARGINGS AVINDAARBA PINGINGERING CINCINGGGG AITGINGGGGT
45	TTCTAGCATG TGATTACTTC ACTCCTGGAC TGTGACTTTC AGTGGGAGAT GGAAGTTTTT	840 45	CONCENSION ECONOCION PRODUCTION PRODUCTION CONCENSION PROPERTY OF PRODUCTION
	CAGAGAACTG AACTGTGGAA AAATGACCTT TCCTTAACTT GAAGCTACTT TIAAAATTTG	006	CALCADATA ON CALCADATA INCOME. STATEMENT CALCADATA CALCA
8	AGGOTOTIGGA CCAAAAGAAG AGGAATATCA GOTTGAAGTC AAGATGACAG ATAAGGTGAG	096	AIMAICLIMA GIICAIGMUU CIITARUUU CUIMAUUUN AACAIMAAIL ALAUUIUIU
2	AGTAATGACT AACTCCAAAG ATGCCTVCAC TGAAGAAAG GCATTTTTAAG ATTTTTTTAAA	50	TAGGGAGGAG KCAAAIGIAG GICAIGITIT IGITGGIACT ITCTGITTIT IGIGACTICA
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;	MILLIOLON GAMBAGINCI ANITITCALI ASCANITANI ABACCIATAC	-	TOTCCCTOTT CATTACATOT CATTCAGTAG GTGGGTAGCC CTGATGGGGG TCGCTCTGTC
ç	ATGCAGAAAT GAATACAACA GAACACIGCT CITITITGAIT ITAITIIGIAC ITTITIGGCCT	1140	TOGAGCATAA CCCACAGGGG TITTITICTGC CACCCATCC CTGCATGCCT GATCCCCAGT
	GGGATATGGG TTTTAAATGG ACATTGTCTG TACCAGCTTC ATTAAAATAA ACAATA	1196	TOCTATACCC TACCCCTGAC CTATTGAGCA GCCTCTGAAG AGCCATAGGG CCCCCACCTT
9		09	TACTCACACC CITAGAATTC TGGGAGCCAG TCTGCCATGC CAGGAGTCAC TGGACATGTT

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5 S AAAAGGTATA TATGCATATA TCTATATATA ATATGACGCA GAAATAAATC T GAGATGATTC TTTCTTOGCC CTGGCCATCT CGGGAAGCTT GATGGCAATC CTGGAAGGGT GOGAATOCTG CTGCTTCAAC CCCAGAGCCT AAGAATOGCA GCCGTTTCTT AACATGTTGA CATCCTAGAA TCCTGTCACA CTACAGTCAT TTCTTTTCCT CTCTCTGGCC CTTGGGTCCT TTAATCTCCT TTTGTGAGTT TGGTGGGGAA GGGAAGGGTA TATAGATTGT ATTAAAAAA 1740 1680 1620 1560 1791

(2) INFORMATION FOR SEQ ID NO: 224:

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E SEQUENCE CHARACTERISTICS: € LENGTH: 2517 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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Ĕ. SEQUENCE DESCRIPTION: SEQ ID NO: 224:

ACACTAGTGG ATCCAAAGAA TTCGGCACAG CGGCACAGCA TTGTTGAGCT TTTCTGTGTG

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AAAAGGACGA TGCCATCACC GCATAYAAGA AGTACAACAA CCGGTGTCTG GACGGGCAGC TOTOGOCCCC TCAACCGAGC TCGACTOGTC CATCCTGGGG TAGCGASGTG GIGTTTGTGA 180 120

30

TOCGGCTGAG TGACAGCCCA TCAATGAAAA AGGAGAGCGA GCTGCCTCGC AGGGTGAACT CGATGAAGTG CAACCTTCAC ATGAATGGGA ATGTTATCAC CTCAGACCAG CCCATCCTGC 300 240

CIRCCICCIC CICCAACCCC CCIGCCGAAG IGGACCCIGA CACCAICCIG AAGGCACICI 420 360

35

TCAAGTGCTC AGGGGCCTCT KTGAGCACGC AGCCCACAGA WITCAAAATC AAGCTTTGAG

CAGGGGAGTR AGGCAGCCAG AAGTGGGGGC AGAGGAGGGT GGCTCTGTTT CCCCAAGGCA

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GGGCTCACTG

CITICACIGI

GCTTAGGGTA GGGATGGTAA

ATATCCTCCC

TIGCATIGGCTT

2160 2100

CICATCCCAA AGCAGGIAIC TICIGGITGI CACAGAGITT CATIGAGICC

ACGIGGCCAT CIGGAGCIGG TGCTATAGGT GACCATCIGG TACATIGAGG

GATAAAGAGC TICCICACIG GAIGGGACCC GCCITICIGI GIIGIGIICI GCCCIGIGCI AAGCTIATGA CCAAIGGGCC AICGGACIGG AGACCCCIGA TIGIGGGAAG GGTIGCCAGG

CGTTAACGTT TCCTGTAGTA TGTTTCTTCA TCTCATCGCC AAGGTAGGCT

TOTOTTTTTM AGTGTGTGCC TCCCCGAGCC TCAGCCCCAA GCTGATTTCT TATCTGGAAA

TOGTACACTO AATTCTCTGG GTGGCTTTCT TOTGGCCCCA TGGGATGCAG CGTGGGGGCT

GICTGAAGGA CCCIGCITIT ICCAGGGGCC GAGGGGCTOC CITICCITIG IGTGIATIAA

GCTTTTCAAA CAATGGAGGG GATGGAGAGC CCTGGTGTCC TGACGGAGC CAGGTCGGCC

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TGAGAGCTGT

GCCGCTCCTC

TOTOTTOTCA GIGGAGGIGG CIGGGIGGG AGCAGGICTC

960

900

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CIGCACCGCT GGGAAGAGPC TATCIDAGCT CTTGGCTTGG AGTCCCGTGT CGTCTCCRCC

1080 1020

AGGCCTCTTG TCCTCTCCCC AGTGGCTCCA GGCCTCACTA GTGGCAAGGG CAGGATGAGG

840 780 720

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660 600 540 480

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CCTGTGCAGA AGGAGAGGAA GGGGCATTAA GAGATGAAGG

GIGATTATGT

ATTACTTATO ATCIGIGICI AGGAGAAGGT

2460 2400 2340 2280 2220

2517

TCCTGTGTCC

TECTECACIT CECATGEETC TATGITACEC

GCCTCCTCCA CTCTATAAGC AGTCATCTTG

GGAGACCGGG

CATTICIGAA TAAACATITG TIATICCINA AAAAAAAAAAA AAAAAACICG AGGGGG

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30 23 cregregate

ACCCAGCITC AGIGIGCAGG CICIGAGGCI GCCCAGGACG GGAAAGICCA AGGAAGGGGC TCCACTTGCA GITCITTAAA GAATGCIGCT

CTGGCCCAAG AGAGCATCCA GAAGGCAGTA GGACCTGGTT CAGACTTOTO GITTAGCAGCT GGAAGACATT COTOCCACAC TITTOCCTTO TITCAGGTAC TGGGAGCCGG TITTATICIC CTAACCCTIT

2040

1980 1920

AAACCAGAGA CTCATGTTTC CAGGGGTCAG TCTGTCAGGC AGGAAGGACC CAGGATTTGA GOCACATGCC GAAGATACTC AAGAGCTCCC AAGATTTGCT TGAGGCTAGC CCAGTGAAAA 1860

GAGCCAGTTO GGGAGCACGG TICTGGGAGC TCTGCAAAAT CAGTAGCAAG TGCTGGAAAA 1800

20

GITGICACIG AGIGGICCCC IGCIGGITGG GAGIGAAGAG AATCCAGGCI GGCAGAGCIG 1740

1680 1620 1560

GOCCCAGACA GITTOGGACGA AACITTCAGAG CCCAGGCAGT CCCTGAATGA CCAGGCCAGT AGAGGAGGAG GAGTCAAGGG AGCAGGGCAG CTCTACCAGG CAAGGTGTTT CCCCCAGCATA TOCCACTIAT GIGAGIGACC CCATCCATCC ATGACCAGAG GATTATITIC CIGCCITGGC 1500

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GACAGIGGIC AGGGAIGCCI GGAGGCATAT ATCCAGCIGC CACCAAGGGG CACIGITITGI 1440

TITIGAATOGA AGAGGICAGT TIGTICCIGG CICICCAITI CIGGCCICAG TIGTCIACAG CCTYTICAAAG GCTCAGCCTC CCATTOTOCA GTGCTTGGGT TTGGAGCTTA 1380 1320

5

CAGAGGAAGT TCTCCAGAGT TCACCTTTCC CTTTTCCTTG AGTIGTGCTG AATGCCCCAC

TCTGGTTGGG COGITICICAC CICGCCIGGC ACITAACCAC ACCCIGGITT IGIGIAGCCG CCAGCICICI CCCAGCICIC TITICCCTICI GGGIGICITI GCIGGGAGGG GGCIGIGITG IGAGCCCICC 1260 1200

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5 INFORMATION FOR SEQ ID NO: 225: Ξ

SEQUENCE CHARACTERISTICS: (A) LENGTH: 2424 base pairs STRANDEDNESS: double TYPE: nucleic acid

SS

TOPOLOGY: linear

452

(X SEQUENCE DESCRIPTION: SEQ ID NO: 225:

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	TIGIANCIAA TOGAGGATIG ATTOTAATGA CAGAGIOTITI CAACACITIG CACATGAIGT	09	CCCCCATGCT GTTTGTATAA GTTTTGCTTA TTTGTTTTTG TGCTTCAGTT TGTCCAGT
•	ATCACGAAGC TACAGCTTGC CATGTGACTG GAGATTTAGT AGAACTTCTG TCAATATTTC	120	TCTCTGCTTG ANTGCCAGA TAGATTTATA GCCTTAATTC TTGGTCAGGC AGAACTCC
י	TITICOCITITI GAACICTACA COCCCITATC TICAGAGAAA AGAIGIGAAA CAAGCAITAA		atgaaaaaa cttgcatctt cagtatactt cctaaaaggg aatcagataa tggaatatg
	TCCAGTGGCA GGAGGGAAIT GAAITTGGCC ATAAACTGTT AACTCTTCTT AAITCCTAIA	240	TTATGTAATT AAGAGTTCAC TTTAGTGGCT TTCATTTAAT ATGGCTGTCT GGGAAGAA
0	STCCTCCAGA ACTINGAAAT GCCTGTATAG ATSTCCTCAA GGAACTIGTA CTTTTGAGTC	01	GGGTTGCCTA GCCCTGTACA ATGTAATTTA AACTTACAGC ATTTTTACTG TGTATGAT
	OCCATGATIT TYTICATACT CTGGTTCCCT TICTACAACA CAACCATTGT ACTITACCATC	360	GOTOTOCTOT GTOCCAGTIT TOTACCTTAT AGAGGCAGAT TGCCTCCGAT CGCTGTGG
<u> </u>	ACAGTARIAT ACCARTGTCT CITGGACCTT ATTTCCCTTG TCHAGAIART ATCAAGCTAA	420	CITATTATCA AAATTAAGIT TACTIGIATA CGGAACAACC ACAAGAAATT TGATTCTG
2	TAGGAGGGA AAGCAATATT CGGCCTCCGC GCCCTGAACT CAATATGTGC CTCTTGCCCA	480	AAGANTCCTC TTTAGCTGTG GCCTGGCAGT ATATAAATGG TGCTTTATTT AACAGAATA
	CAATGGTGGA AACCAGTAAG GGCAAAGATG ACGTTTATGA TCGTATGCTG CTAGACTACT	540	CTOTOGAGGA AATAAAGCAC ACTTGATGTA AAAATAATTG TTTTATTTT ATTGACATC
20	TCTTTTCTTA TCATCAGTTC ATCCATCTAT TATGCCGAGT TGCAATCAAC TGTGAAAAAT	600 20	CTGATTGATT GCTATTCTGT GCACTNAATT AAACTGATTG TGATGACTTA AAAAAAAA
	TTACTGAAAC ATTAGTTAAG CTGAGTGTCC TAGTTGCCTA TGAAGGTTTG CCACTTCATC	099	adaladada adabadada adaa
3,5	TIGCACTOTT CCCCAAACTT TGCACTGAGC TATGCCAGAC TCAGTCTGCT ATGTCAAAAA	720 25	
3	ACTIGCATCIA GCTITITIOTOT GAGGATCCTG TITITOGCAGA ATAITATAAA TOTATICCTAA	780	. 300 . ON CT DES EAST HOUSE IN 10.
	TGGATGAAAG AACTITITITA AACAACAACA TITOTCTACAC GITCATGACA CAITITCCTIC	840	10) THE CHARLES CONTROL TO THE CONTROL OF THE CONTR
30	TAAAGSTICA AAGTCAAGIG TITIICIGAAG CAAACIGIGC CAATITIGAIC AGCACICITA	30	(1) SEQUENCE CHANGITERISTICS: (A) LENGTH 1080 base pairs (n) more
:	THACAAACTT GATAAQCCAG TATCAGAACC TACAGTCTGA TITCTCCAAC CGAGTTGAAA	096	
ž	TITICCANAGE NAGIGETIET TIMANIEGGG ACCIGAGGG ACTEGETITG CICCIGICAG	1020	(U) IOPOLACE: LINGAL
ર	TACACACTCC CAVACAGITA AACCCAGCTC TAATTCCAAC TCTGCAAGAG CITTITAAGCA	1080	(XI) SEQUENCE DESCRIPTION: SEQ ID NO: 220:
	ANTIGCAGAC TIGTCTGCAA CAGAGAACT CACTCCAAGA GCAAGAAGCC AAAGAAAGAA	1140	ATATAGACG GATAATCIGI TTACATICIG TICITCICGA TOCACICACA NGCGGGIA
9	AAACTAAAGA IIGAIIGAAGGA GGAACTIOGA IIPAAAAGGGG GACAIGIIIAGG AGAIGAAGA	40	TAGGTGACAA GAAAACAAAG ATCTTATTCA AAAGAGGTCT TACAGCAACC CAACGTCTC
<b>!</b>			TUTTUCCATA GTAAAGATGA CGGCGCTTG AGGTAAGCTA CAGGCAACAC CACTTCCGC
	ACCACACTOT ACACAGCTOC ATCAGTGACA TGAAAACAGA AACCAGGGAG GTCCTGACCC	1260	TITICITETIOC OCCCTOOTICC AMONTOOCOG ATGAAGCCAC GCGACGTOTT GTGTCTGAG
45	CAAGAGCAC TICTGACAAT GAGACCAGAG ACTCCTCAAT TATTGATCCA GGAACTGAGC	1320 45	TCCCGGTGCT GAAGACTAAC GCCGGACCCC GAGATCGTGA GTTGTGGGTG CAGCGACTG
	AAGATETTEE TTEECETGAA AATAGTEETG TTAAAGAATA CEGAATGGAA GITECATETT	1380	AGGAGGAATA TCAGTCCCTT ATCCGGTATG TGGAGAACAA CAAGAATGCT GACAACGAT
	COTTITICAGA AGACATOTICA AAFATCAGOT CACACCATIGC AGAAGAACAG TCCAACAATG	1440	GOTTCCGACT GGAGTCCAAC AAGGAAGGAA CTCGGTGGTT TGGAAAATGC TGGTATATC
S	GTAGATATGA CGATTGTAAA GAATTTAAAG ACCTCCACTG TTCCAAGGAT TCTACCCTAG	1500 50	ANCANCING CARABBARANCE TOTOCACANTO ACTUATION TOCTARGES PARTICIONAL
	CCGAGGAAGA ATCTGAGITC CCITCTACTT CTATCTCTGC AGTTCTGTCT GACTTAGCTG	1560	POSSONER SHAMESTAR STANDARD ST
ý	ACTICAGAMO CIGITGALGGC CAMOCITITGC CCICCCAGGA CCCICAGGIT GCITITATCIC	1620 <<	TOO OUT THE TOTAL TOTAL TOTAL OF TOWN TO THE TOTAL TOT
3	TCAGTTGTGG CCATTCCAGA GGACTCTTTA GTCATATGCA GCAACATGAC ATTTTAGATA	1680	COMMITTE CLIMICANT CHITCHEN CLIMACY COMMITTE CLIMACY
	CCCTGTGTAG GACCATTGAA TCTACAATCC ATGTCGTCAC AAGGGATATC TGGCAAAGGA	1740	CALINACITA ICITATOSCI LIGGOCIOS GICCATOSCI CACAGOSTA AICECTON
99	AACCAAGCTG CITCITGACA TTAGGTGTAG CATGTCTACT TITLAAGTCCC TCACCCCCAA	09 0081	TGATTCHGAA GGGCGTCATC CAACACAAAG AGAAATCCAA CCAATGAAGA ATCAAGCCA

CCAGGTGGAG CAACATGCGA TICTGGAGGC ACGGGGGTAA CTGAAAGTGA GTACATATAG GCTGCACAGG GAAGGGAAAG ACTOGGCTTT GGACAATCTA GAGGTAATTT ATATCCGCCC TAGGCATTGC TGGGGAAGAA ACAAACACAC ACCAAACAGT ACTGCTACTT AGTTTCTAAG AACTCATCTA ACTGCTTCCC CGGACACCCT CCACCTCTAG TTGTTACTAA GTAGCTGCAG TEAGGCAGGG CAGAGOGACC TTTGATAGGC TACGATACTA TTTTCCTGTG CATCACACTT 455 1020 960 900 840 780

20 2 INFORMATION FOR SEQ ID NO: 227: Ξ

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TETTTETOGT TICTOGAGAT AACCCATCAA TAAAAGCIGC TICCTCIGG TAAAAAAAA

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OTCCCTOTOT AGCCAG

CICTICITOT TGAGGGCTAT GCCGGGTGGC ATOTTTCCAG GGAATCTGGA AGCGTTTAGA

1336 1320 1260 1200 1140

ATCUACCECT TOOTGETACT TOOCTGTOTO TACAGOODAG GOOTTAAAAA TOOTATTTTTO TICTOTOCIO TIATIGAGAS CACASCOCAS CIGGOCCITO CATITAGOIS GOICAGGGIO ATATATETTE ACTOSOTTEA TOSSCACETE CETTOSTOCE ATECATACAT CCAGOTTGAA

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(A) LENGTH: 1336 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

X. SEQUENCE DESCRIPTION: SEQ ID NO:

TIGOATICAC AATTACTOGG AGGCAGOCAG GOGCAGTIGG ATGCTGGOGG TGGCTGCAIG

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ATOTOMANO GIGGOTOCTT CICTATOTOT COTTICTOGT GGGCAATGAG ATCGCTAAGG AAGATGAGTT TECTTTECAA TOGTTTECCA TETGGECATT CTTECCCAAA GCATAAGTAG GTGAAATCAC CAGAAAACAT CTCGTAGATC ATCCGAGCCA CTACTGGAAT GACCTGAACC TTATCCTTGA TGTCTGGGGT TGAGATCTGC AGGCGGACAC TGCCACTATC AAAGGATCGT GSCTGCCASC TCTCCTGGGT TTGAAGGATG CGGTACASCT GCTTCAGCTG AGCAACGATG

AATGTITCCA GACAAAATAG CITGACCITC TITTGTCTCT CAATCAGGIT GGGAGCAACA

420 360 300 240 180 120

480

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GTGAACAGCT TCTGCTTCTC ATTGGTAATA TGCTCAGGAC CTGGGAATTT GACCTGCTCC ATGGCATGAC CGCTGACCTC CAGGATCAGT CCTCTGTCCA TGACGTCCAG CAGCTTGCTA AGTGATGGGG CACATGGCCC AGACCAGTAC ACCTTGCACT GGCACAGYCT GATGGCATAA

> 600 540

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AGNOTIGAÇOG GACCAMAGAG CITCOTOCIOG ICAGGCATOG GACCCAGGIC COCATAGAAG

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CTGAAACTTG ATGTCCAGGT CAGTCATTGG GAGAGAGCTG ATCCACAGTT CTGGAGAGCT

ATAGAAGGRC TGTATAGGTG CCTGGGGHAC TICCATCICC AGGGGTICAG TITTGGGCCA

AGTEGGCAGC CCTGAGGGTT GCTCACGGTC ATGGTCCTGC CCGTACTCCT

TCCCACGGTA

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AGACCCTOTO GATCCTOSOT TAATGATCGA SCCCTOSOOC TGAGGGATOT CACACACTTG ATCCAGCTCA TETTECTICAT CITETTCATE CACATCATTA TECTTETCAT CECAGGGAGE

> 1080 1020

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ICTIGGICTY GGAACICCCY GGCAFIGGGA ACAGAGCAIT TCCAGCAITT GITGTIGTIG

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CACTRICCTICC, GGSCTGCAGT TGCCCACACT GCAATTGCCC ACACTGGCTG GCGCCATGGG

AUGITCAGGA AGGGGAAGGT GTCCTGGATG GGAACATGGT GCTGCGACTG

960 900 840 780 720 660

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£ SEQUENCE DESCRIPTION: SEQ ID NO: 228: 20

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 2043 base pairs 2

(2) INFORMATION FOR SEQ ID NO: 228:

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SEQUENCE CHARACTERISTICS:

23 GACCTCAGAC CCCACCCACA CTCCAGATCO AGACCCCTOC CTCCCCCCGO CAAATOTCCT TEAGETGGIC CETTECTION GICCIGGGG ACCIGGIGG GGCCICTICC TGGGAGCCAN

ၓ 3CAGAGGCCCA GGGTCCAGCA GCCCGGSGGG AACGGGTGCT GCCTSTTCCT CCAGTTAGCT CICAMPICCE CITECCIDAG CICCITECCA AGGACICCIO GICACIOCCI OCIGIGCAKI COCGETOCET TOCAGECTOC ACTITICANEA TOCTEACECE CAGGAGAGITE CEACTOOCEC 120 S

GETIGGAETIGG TECEGGAGETIG GETTEETTAE CAGAAAAGEE TEAGEETTEE TETIGGAAGEA CICIOCCCIC AGGGICAGEI GCCCAGACIG GGGGGAIGC AGAGAGGCAG GIGGGCIGIG CAGGOCTOGO TOCTGAGCAG CTGGTTTTTCC TOCAGGAAGG TTGGAGCAAG CAAAGTCCTT CCAGYTCAGG TCTGAGACCC GTGYTGAGTA AAGGTCTGAG CAMCGACCGT GCCCTCTGCC 240 540 480 420 360 300 180

INCOCCUTTO TODOCIAMODO GGAAGOGOTIC CITITAAGOGG TOTOCITITOC CAGTIGGGGAG 660 600

CAGICIOSCC CIGCCCCTA CIAAAGCCIC ISCICICAGC ACTITICCCCC AAGICCIIGI CCAGGCTGGG AGGGTTCCTT CCCTAGCTCC CCATCTGCCC CCGCTGGTGA GAGTTGGGCT NGCCCAGCTG GGGGGCCGGG ATGGGGGCTT CTCTCTCTGG GAGGGGTGCA GGTGCCCTCC CCTOGTGAGG GGAGGTOCTG CTTTTCTOCC CCACCTOCCG OCTOGTTCCA GCAGCOCTOG TITITICATIT TITICITICCG TCTTTCTTCT TGAGTTCACG GITCAATATI GCCTCCTCGC TITAAATTIC ACICATITIG TATAAACCCA GCAGGCIGGI GITIACTIAG CCCIGTAGCI AACTIGETIG AAGGIGGGIT CIGGEIGEEA GEEAGIEEET GGACAAACIE TECIGEEEET 1020

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à	1717 a	THISTORIAAAA	S OW	WO 98790448	PCT/US98/04493	
	457			458		
	TITITACTENE CTANCECTIN GANANTGANI GTINGANGST GECTGCCGAG GEGGGAGA	1140	•	PARTICIPATE DESCRIPTION OF THE PROPERTY OF THE	. 087	
	STOTTTOCTC GOOCTOGAGA AGGCTCTGCT CAGCCCTGAG AGTCCCTTCC TGCCCCACGG	1200		TUTIONIONE CENTROLLING CENTROL STREETS OF TELEGOOM STREETS INTROLLING	, to	
5	ATACTOSCAC TTTAAAAAGS AACCTGACCG CACACTGTCC AGACGAATTG GCCCCCAGAA	1260	۰ ۲۰	ביונסניות שייניינינין ואפריניאינט מששטטעון אינייניאין אינייניאיני	<u>}</u>	
	GATGGGGAGT TCTGTCCTGC CCTTCTGTGT CTGCGTGACC TCACCCAGCC TAGGAGGGAG	1320				
9	GTOCATTCAG GGTAGATTTG CCTCTCATTC AAAGTTCTGG GGCTTTGGGY GGAAAACAGC	1380	0	(2) INFORMATION FOR SEQ ID NO: 230:		
2	CAGCTITIGGC GCTGTTGGGG AGACTCCTCC AGACCAGGAA CCCCAGAAGG AGACAGAGCC	1440	2	(i) SEQUENCE CHARACTERISTICS:	-	
	TGCCACATCC TCCCACGCCA GGCCCTGGGC CAGGGTGATT GGACTGAGAA TTTGGCCACA	1500				
15	ACCADANTIGA TOCTGOCTGG AACCAGAGGC CAGAAAGCCT GGCCTTGTCC CCATGTGGGA	1560	15			
	GCOCTIOTICET CAGCCCTCTT GTCCCCTTGA GCTCAGTGAA TTCCCACCAG GTGCCCACAG	1620		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 230:		
	CTCCTGGACT TCAAATTCTA TATATTGAGA GAGTTGGAGA GTATATCAGA GATATTTTG	1680	7 02	AATTGTGAAA TATTAGAATA TTGTTACTAT TTGACCCAAC TCAAAATCTC CATGGGAAAA	09	
3	GANAGGACIT GOTCTATGCA ATGTCACTIT GGAATCTICT TGAAAGTITA ATGTTTTTAT	1740		TACCTOTCGA TACCCACAGT ATTGTTGAAA ATAATCAGAT GCAGTATCAC AGCTGTGTCA	120	
	TAGGAGATIT AAAGAAAATA AAGGICTACA ATATCTITAG GITITITITI TITCCTGTIT	1800	Ü	GACTETAGTA CCAGTTGGGC AATCAAGGCA CAGCTAAAAA TTGAAAACAA AGATCTGGAC	180	
25	ACCICIANA CTGACCACAT GOCATOTCTA TCAGGATGGA GOCTOTCCAT STICTCCTCT	1860	25	AACAAAACAG CCAAAGGTGG GGGTCAAGAA GCTCTGACGT GTACCTAGCT GTAGAATGCT	240	
	GICTITAGGS AGGIGATAAG GAGATGSSCS RAGGGGTGTT TITTITCTITG ACTCCCTCC	1920	_	ATGCACACGT GCCAGGTGTA CTGTGCATAT CCAGGAAAA CTGCAGAGAG CCCCAGTCTT	300	
ç	TITCTAACAG AATOTTOCCA CCACTOCITG AGTOGOCIOT GITTGTTOCT CTOTCCCAGC		ς.	CANCTETGGT TEACEATEAG CTETGTGTAA GCAGGAAGTG AAGGCTAAGG CAGATTTAAG	360	
2	ITCTOTICIA GAAAIAACA TIOTIAGGGG AACTCAGGCT AGIGTCAGCG TCTTGGTTIG	2040		CTCTGAAAGC ATTCCACAAC ATACACACAA ATCGTGCAAA GCATTAAGGA AATCTTGTTÄ	420	
	500	2043	Ü	CTGCTAAGTG TTGCTGACCC AGGACAA	448	
35			35			
•	(2) INFORMATION FOR SEQ ID NO: 229:			(2) INPORMATION FOR SEQ ID NO: 231:		
4	(i) SEQUENCE CHARACTERISTICS:		40			
	(A) LENGTH: 540 base pairs (B) TYPE: nucleic acid (C) STRANDERFES. Achid			(A) LEWGIN: 40' Dasse pales (B) Type: nucleic acid (C) STRANDERMESS: double		
45			45	(D) TOPOLOGY: linear		
}	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 229:		}	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 231:		
	TAAAAAGAAG CGGGAGAATC TGGGCGTCGC TCTAGAGATC GATGGGCTAG AGGAGAAGCT	09	Ü	GRATGCTGCC CCAAACCAAT ATGTGTGGCT GCCTTTWACC TGACTTCTCC AACATGTAGC	09	
20	STCCCAGTOT COGNOAGACC TOGNOGCOST GAACTCCAGA CTCCACAGOC GGGAGCTGAG	120	20	CCCAAGAGGA GGCCTCTAGA CTRAGGGAGG GGCTGGTGAC CCAGGTGTGG TGGGGCTGCA	120	
	CCCAGAGGCC AGGAGGTCCC TOGAGAAGGA GAAAAACAGC CTAATGAACA AAGCCTCCAA	180	-	TGARACTACC AGAGAGACAG ACATTCTGGA ACTCACCCTG GGGGATCCAG TGGATCTGCC	180	
8	CTACGAGANG GANCTCAAGT TTCTTCGGCA AGAGAACCGG AAGAACATGC TGCTCTCTGT	240	۶,	TATIGGTCTGG TOCACCCCAG ACCTOTGAGA TOTTCCTCAT GAGGATGCAC TYGTGCTTCT	. 540	
3	GGCCATCTTT ATCCTCCTGA CGCTCGTCTA TGCCTACTGG ACCATGTGAG CCTGGCACTT	300		GCAAGTATTG CTGCAGCTTC ATAGTGACTC CCACCAGCAC CAGCAATACA GYTAGCTACC	300	
	CCCCAACACC AGCACAGGCT TCCACTTGGC CCCTTGGTCA GGATCAAGCA GGCACTTCAA	360	μ.	TOTOGCCTTG GATCTCAGCC AGCATGGCTG GGAGAGGGAG CARCTGGGCA TGTACCCTAA	360	
9	GCCTCAATAG GACCAAGOTG CTGOGGTOTT CCCCTCCCAA CCTAGTGTTC AAGCATGGCT	420	<b>*</b> 96;	ATOCTIOTTAC CAOSGAAGGA CTCCCAGAGT GAAGACAAGT AGGGACT	407	

. **U**s

(2) INFORMATION FOR SEQ ID NO: 232:

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25 20 5 2 TACATTITICT CATTAGOGIT RIGATOCTICA GIATCITICC AAGTOCCAGG CAGROCTING AATTIGACAA AGGGIGICAT AIGCTITICCT AACCIGAWIT GIATIAACAT ICACAGAGCC GGAAAATTGA TACTTTTAAA GCATATTCTT CTATGAGCAC TITITIGAGCA AGAATIGCCAG AAATAGCCTT CATTICTACC CTGCAAAATA AICCAGATCT AGGITICCIA AACIATAAAA GCAGATITIG CITITIGITIG TIAAICATAG GCATGGCCGA TCCCAATTTA AAAATGCAGA CITITICIDAT CAMACATACC ATTITITIOTA TITICACAACT ATAGACAGIC ACTICIOCAG GCTGTGTATT GTGACTATCA GCATTCTGGT GCAAATGAAC TECTGARTIG GARGAGGAAG RACTICIGIT TACAGAAAAC YGTATIGITA TATATGICAG GCTTTCTAAA ATGRANTCAG TTTCTAAAGT GAAACATGCA ATATTTATGC CCCACGTTTC TAATTTTGGAG CAAATCTAAA AG TCACTATCAA GAGCCTGCAG AGCCATTTTC CAGACCTGTG ATTGCCCAGA ACACATAGTC GCATTOTOGA TTAGCCTGAG GCTTAAAATC AGATGCATGT CTOGTAAGAT GACCACTGTC ACTOCTITAT CCAAGAATGC TGAAAAATAC TGTTCTATCC AGGICCICCY AGTGAAACTT TITTCTCCAT CATCGACTGT TCTGACTGAC 900 660 8 540 480 420 360 932 840 780 720 300

2 INFORMATION FOR SEQ ID NO: 234:

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E SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (A) LENGTH: 2786 base pairs STRANDEDNESS: double

(D) TOPOLOGY: linear

35

S 5 8 S TTAGCAGGGT GAGCTGTTAA AACAGCACAC ATCTCTCATC CCCTCTTCCT TTATTCCCCC TEAGTECCCC TOCCAACCCT CCATATOGCT CTCAATOGTG CTCACTTGCT TGGAAGCAGG CTGGGTTTCA GAAAGGAAGG ATATATGGGG ACCACCTCCC CCTTCTTTGA TCCCAGCATC CAAAATCACT ACAATAGCCT AGIGCITTIT TOGAAGCCTT TITAGGGAAG AATGITAGGI CATGGGATCC CTTCCATAAC AGGTACTTTG AAGGCAAGAC ATAGGGTTGA AGAAGCACA TOTTTTTOTO TIGIGGAACC TGAGATICCT TATTTATTAA CAGGAAGICT GATTITITIT TCAGTGAGAC GATGAGAAAA GTCCCAGGCT AATGGCAGAA ATTTGCACTT TGAACATGTG CICCCAATAG GGAGGGGSCT GCCCTCTACA GTCTCTTTGA CTGTAAGACA GGGCTCTGTA CCAGCCTCTG MAATCATAGC TCTCCAGTGG CTTTTAAAGA AAGCTGGTCC TCAGCACTA TTTTGGAGTC TTTGTTGCTA TATTTTGTGG GGCTGGGAGA GAGAGATTAG ATTATTTTGA (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 234: 240 180 120 540 420 360 300 600 480 8

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	461			462 ,		
ą	TCATOGTAAC TAGTATGCTC TITGAGATIT TTACAGTGTT GAAACTIAAG AATTITGAGA	099		ACTAGCATCC ACATAGAGCA CITGAACCTC CITTGTACCT GITTGGGGAA AAAGTATAAT	2460	
g	OGOTGAGGAG GGTTGTTCAG AATCTAAATT ACAGATAGAT GATTGTTTCT TGTGAATTTG	720		GAGISTACTA CCAAICTAAC TAAGATTATT ATAGICIGGT TGTTTGAAAT ACCATTTTTT	2520	
5 111	FITCHTITIC TITITITIES ECCLEDICAL TECCHIACAT FECCETIOS GOCCATOLET	780	S	TCTCCTTTTG TGTTTTTCCC ACTITCCAAT GTACTCAAGA AAATTGAACA AATGTAATGG	2580	
ğ	GOCICCTICC TITITIOTITIC TITICITICS TIANCAGITIC AITICEACTIC CCUSTIAGIS	840		AICAAITTIAA AAIAITTIAI TICTIAAAAG CCTTITITIOC CTGTTGTAAT GTGCAGGACC	2640	
	ARGACACTO CTOTTAOTGA AGGAACAAG TCTATGAGTC CTAAAATTT AAGTCAAAGA	, 006	9	CITCICCITI CAIGGGAGAG ACAGGIAGIT ACCIGAATAL AGGITGAAAA GGITAIGIAA	2700	
- } ⊃	AACTIGCTCT GTTTCCCCTT TAGTAACACT TCTCAAGAGG AAAACTTCA ATAGCCAAAG	096	2	AAAGAAITA TAATAAAAGG GATACTTIGC TTTTCAAATC TTTGTTTTCT CTTATTCTAG	2760	
£	THANDANC DAINDANA TRECTINGE TITCACCIAA MITCHGGG ANCACAATIT	1020		GTAAGGCATA TTAAAAATAA ATATGT	2786	
5 00	CCTTGGGATA GAGGTTGTGT TGGGGAATAG ATTGCTTATT CCTGTTCACT GGAGAGAAA	1080	15			
ğ	GENACIOITI TIGNACAGO TCANACCOCC AGAAGCCCCA AATCCTATIT TGGCTCATCT	1140			٠	
	TCAGGIAAAG AGTAATTCCT ATCCTGTGTG CCTCAGAAGC TAGAATGGAA GGCTTACCCT	1200	ç	(1) INFORMATION FOR SEQ ID NO: 233:		
0. E#	ATTCATTGTT TATTGTCAGA AATGCATGAT GGCTCTTGGA JAGAATGAGG TTTTTGCTGGA	1260	2		•	
\$	aaaaaaaa agaacaottt gtotttcaca aacatgoctt atcaattttt tcaaagaatt	1320		(b) Tire: nucleac actu (c) STRANDELISS duble (n) Thomator lines		
Si Fi	CTTTTTCCC AAAAGAGSA GTAACAAAT GTCATTTCTG AAAGAGGCTT ACTTTATACC	1380	25			
Ž	AACTAGTOTO AGCATTTGGG ATGCCAGGGA ACAGAGAGTG AGACACCTAC AATCACCAGT	1440		TO THE PROPERTY OF THE PROPERT	. 09	
	CICAAAIGCG CIAITGIFFIC TITTCAGAGT GITGCAGAIT TGCCAITTICT CCATAATATG	1500	9	CONTROL STICOGRAP ADMINISTRAL CONTROL	120	
8 >	GGGATAGAĄ ATGGAATAAA GATAGAAGSG ATGTAGAATA TGCTTTCCTG CCAACATGST	1560	3	בייניים ביינים ביינים ביינים בי		
Ę	TIGGAGICGA CITIGGIATA ITGACIAGAI ITGAAAAIAC AAGAITGAIT AGAIGAAICI	1620		TACCTGCATA ATCACAGCTA TGCATCTATT CAAGTGATG ATCTGTGGGA TAGTTTTAAT		
ν. Š	ACAAAAAGT TGTCCTCCTC TCAGGTCCCT TTTACACTTT TTGACTAACT AGCATCTATA	1680	35	GAGGICACAA ACCAAACACT AGTIGTAAAG AGAATGATGA AAACCTGGAC CCTGCAGAAA	240	
Ę	TICCACACIT AGCITITITIG ICACACITAT CCITIGNCIC CGRAAATITC AITITGCAGTG	1740		GONTITICAT INSTRACTION TOWNSHAME CONTINUED INTITIATION CONTINUED C	2 5	
	OTTAGECATE AGAINITITA OCCACETACA CANAACCANA CTGCNITITT ANAAATETIT	1800	\$	TICTITIAN AIRIGANACC TOMMITCAG COTTOMARIA CONSISTACT COCCIOTATO		
E	CHSAGATGGG AGAAAATGTA TTCTCCTTTC CTATACCGCT CTCCCAACAA AAAAACAACT	1860	7	THE CALL AND A STANDARD CALLETTERS INCOMMENTED THE CALLETTERS OF T	, d	
ĄĠ	AGTINGTICT ACTANITIASA AACTIGCTOF ACTITITICIT TICTITIAGO GOTCAAGGAC	1920		ANGITUTION UNANCUTANE ANGALITATEC ELICEIAN	Ş	
55	CCPCITIADA GCIACCAITT GCCIACAATA AATTATIGCA GCAGITIGCA ATACTAAAAF	1980	45			
AT	aittitiata gacittatat tittoctitt gataaagaga tgctgcatag tagagitggt	2040		(2) INFORMATION FOR SEQ ID NO: 236:		
8	GRANTIAAAC TATCTCAGCC GTTCCCTGC TTTCCCTTCT GCTCCATATG CCTCATTGTC	2100	9	(1) SEQUENCE CHARACTERISTICS:		
	CTICCAGGGA GCICTITIAA ICITAAAGIT CIACATITCA IGCICTIAGT. CAAAITCIGI	2160	3			
¥.	TACCTITITIA ATAACTCTIC CCACTGCATA TITCCATCIT GAATTGGTGG TICTAAAITC	2220		(D) TOPOLOGY: 11near		
δ. 1c	TGAAACTGTA GTTGAGATAC AGCTATTTAA TATTTCTGGG AGATGTGCAT CCCTCTTCTT	2280	55	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 236:		
KG	KOTGOTTGCC CAAGOTTGTT TTGCOTAACT GAGACTCCTT GATATGCTTC AGAGAATTTA	2340		AGGATGAAGA GGAAATTATC TCTTGGAITG CTCTCCAGGA AATCCTTCTC TATACTTAA	09	
8	GGCAAACACT GGCCATGGCC GTGGGAGTAC TGGGAGTAAA ATAAAAATAT CGAGGTATAG	2400	Ģ	AAGCICTIGT TCTTTTCTAG GARTCCAATG TGCTGATTGC TGCTAACAGT CAGGGTACAA	120	

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TTRAGGTOCT AGAATTGGTA TGAAGGGTTA ACTCAAGTCA AATTGTACTT GATCCTGCTG

5 S AUGGACATIG CIGCICTIGG TGGIGTIATC TAATTITITGT GATAGGGAAA CAAATICTIT CARAGICATE AIGGGITTIG GARTIGITIT GARTATITITI TETITITITE TIKICCETEC AAATACATOT GCAGOTGACA ATGAGAGARG AAACAGAAAA TOTOATGTGA TOTOTOTOCO TITITOTCAA ATGCNGCAAG AGATAACICT TITTANGAAG TAGCATATGT GAACTATAAT TGAATAAAA TAAATAACWA AACAATAAAA GTTTATTGAG CCACAGTTGA GCTTGGAAAG TITATGAGCC TITIOGGACAT TOGGAATACC CAGCCAACTC TOCACCATCA ATGTAACTCC

> 420 360 300 240 180

7

GGTTTTICCNC ACCTTTTTGG TIGGGC

CCATTACTOC TOTOGACACT CTTOGCTTRG TATWITTAGG GOOGNTCCTT ACCITITITE

1260 1200

ATCACAAARG GCAACCAARG GOCCCCTCTT ARGOCTTTGA GGATTAAAAC TAGTCTTTAT AGGGTICCAC TIGGGCCACA GITTITITIGT TAAICAAACA CCACICICIT AAGROGCIGC

S

CAGOGRAFICT ARAGAGETIST STERRICTISTS TREATRICAGA GRETTRECTISA GRARAGISTICA

1140

1080 1020

aaaaataaaa agacagcaat gactitatat ccaagaaagg aatgtgaatg agtcacttaa

480

5 GTAACAGTGA ATAATTTGTA AAGTTCGTAT TICCCAACCT CITTGGGAAT T

> 591 540

5

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INFORMATION FOR SEQ ID NO: 238:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 734 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

3 INFORMATION FOR SEQ ID NO: 237: Ε SEQUENCE CHARACTERISTICS:

20

X. SEQUENCE DESCRIPTION: SEQ ID NO:

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SEQUENCE DESCRIPTION: SEQ ID NO: 238:

ATOGCAGOGO AGAAGGACCA GCAGAAAGAT GCCGAGGCGG AAGGGCTGAG CGGCACGACC

60

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TOTTTTTANG GTACAGCAGG GAAGAACTGG AAACTCAGAG AAAGAAACTG CCCTTCCATC TACAAAACCT GACTTTACTT CTCCTCCTTC TTTGTTCAAG ACTGGGCTTC CACCGAGCAG

GAGATTACCT GGGGCAATTG ATGITATCGG TCAGACTATA ACTATCAGCC GAGTAGAAGG

CAGGCGACGG GCAAATGAGA ACAGCAACAT ACAGGTCCTT TCTGAAAGAT CTGCTACTGA AGRAGACAAC AATTITAGCA AACCACCTCC GTTTTTCCCT CCAGGAGCTC CTCCCACTCA 300 240 180

CCTICCACCT CCICCATTIC TICCACCICC ICCGACTOTC AGCACTOCIC CACCICIGAT TECHCEACEG GGTTTTCCTC CTCCACCAGG CGCTCCACCT CCATCTCTTA TACCAACAAT 480 420 360

TOCCATCAAG AACCATACAG ATACAGGGAA TATOCAGAAA GAGGTTATGA GCGTCACAGA AGAAAGTGGA CATTCCTCTG GTTATGATAG TSGTTCTGCA CGTGCATTTC CATATGGCAA 540 600

AGACATAAGT CTTCTCGAAG TAATAGTAGA CGTCGCCATG AAAGTGAAGA AGGAGATAGT

8

CACAGGAGAC ACAAACACAA AAAATCTAAA AGAAGCAAAG AAGGAAAAGA AGCGGGCAGT

GAGCCTGCCC CTGAACAGGA GAGCACCGAA GCTACACCTG CAGAATAGGC ATGGTTTTGG

720 660

ATATTAGTAC CAGAAGTAGA TACTATAAAT CITGITATIT TICIGGATAA

SS

TCCAAAATAA AAGAGTGAAT TTTTCATGIT AAGTTAAAAA TCTTIGICIT GTACTATTTC TOTTUAGAA ATTIACCITA AARCTIGITC TOTTIGITAG TAIGAAAAGT TAACTITITT

> 960 900 840 780

60

GCAAGTCGAG AAAANGAAGA ACGACATAGA GAAAGACGAC ACAGGGAGAA AGAGGAAAACC

3

35 30 25 6 (A) LENGTH: 1286 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

120 69

30 6 ઝ GIGITICCIGG GCCTCATCCT GIACIGIGIG GIGACGICCC CTATGITICCT GGIGGCICIG CIRCIBECTA ABETRATTICE CIRCOGRIGGA GOCCOGGAGOT GOCTOGAGOG GCGCCCCCCCC GCTGTCTTTT TCGGCGCCTG TTAACATTCT CTATCTGCGC ACCTTGGAGT CCAAGCTTGT GEAGAGETOT GECAGEGEET EGTALGEAAC GTGGAGTAET ACCAGAGEAA CTATGTGTTE ACCATCCOOC CCTOGAGCAC CTTCGTGGAC CAGCAGCGCT TCTCACGGCC CCGCAACCTG

CCTCTTTGGC CGAAAGGTGA GCCCAGCGCA TCATATGCTC TGGCTGGAGG CATCTCCTTC GIGGTEATEG GETECEAEGE TGECTTECAE CAGATIGAGG CIGIGGAEGG GGAGGAGCIG ecementer acerdacias recosacies occarenter assisciosa asceaecepa 540 80 420

360

240 180 120

300

CAGATOGAAC CCGTGTGAGG TGTCTTCTGG GACCTGCCGG CCTCCCGGGC CAGCTGCCCC CAAGCCCOOG GAGGGATICCC GCCTTTGAAA ATAAAGCTIGT TATIGGGTIGTC ATTICAAAAAA ACCCCTOCCC ATOCCTOTICC TOCACOGTCT GCTGCTCGGG CCCACAGCGC CGTCCCATCA 720 99 90

2 INFORMATION FOR SEQ ID NO: 239:

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734

3

SEQUENCE CHARACTERISTICS:

E (A) LENGTH: 809 base pairs (B) TYPE: nucleic acid

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466

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480 540 999 720 780 840 900

909

465

300 360 420 9 120 180 240 780 540 9 99 720 809 120 180 240 300 360 420 480 COCOCTICC ARCOTOCIANG GGGAACCCGA AGTCCGAGGA GCCCGARGTC CCGAACCAGG CAGGGCAAGC TCCCCGTCTC CGGGCCACT TCCCTCGCCT GTGTTCGGTC CATCCTCTT TETECAGEET CETECECETEG CAGGEGATG ANDCOGAGGA CGGGCCAGTG CETGGCAGCC TCGACCCACG CGTCCGGCAA CATGGCGGCT GCCGTGGTGC AGGGCCGGG CTGAGCGACA TICAGCCCGC TCCTGTCCCC GACATCACGT GTATTCCGCA CGTCCCCTCC GCGCTGTGTG TETACTGAGA CEGESAGECG TEACAGGECC CEGETCCCTT CTCAGTGETG CTCTGTCCTT OCCCOSOGIAG GGATCCCCCC TITICAAAATA AAGCTGTTAT GGGTGTCAIT CAGGAAAAA nementacer accreatace sacreasce rementacer acreasasce accreates TCATCGGCTC CCACGCTGCC TTCCACCAGA TTGAGGCTGT GGAGGGGGAG GAGCTGCAGA TEGAACCOST STGAGSTSTC TTCTGSGACC TGCCGGCCTC CCGGGCCAGC TGCCCCACC CTGCCCATGC CTGTCCTGCA CGGCTCTGCT GCTCGGGCCC ACAGCGCCGT CCCATCACAA COGGOTOTIC AGGOTACCOG GCIGOTIACA GCAGCICTAC CCCICACGAC GCARACAIGG CACCECAGAA GEACCACCAG AAAGATGCCG AGGCGGAAGG GCTGAGCGGC ACGACCCTGC INCESCICITE GACACCTIC GIGGACIAGE AGGECTICIC AGGCCCCGC AACCIGGGAG AGCTOTOCCA GCGCCTCGTA CGCAACOTOG AGTACTACCA GAGCAACTAT GTGTTCGTGT cerescer careerstae rereresta coreectar errections ecretiseers ICITITITICOS COCCIOTIAC ATTOTOTATO TOCOCACOTI GOAGICCAAG CITIGIOCIOT TIGGOCGAGA GSTGAGCCCA GCGCATCAGT ATGCTCTGGC TGGAGGCATC TCCTTCCCCT TOCCGAAGCT GATTCCCTCC GGTGCAGGCC GGGAGTGGCT GGAGGGGGCGC CGGGCGACCA GCAAGTGCAG CGGGCTCCTA CCCGGGTGA GGGGTGGCCT CCGGGTGGGA (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 240: SEQUENCE DESCRIPTION: SEQ ID NO: 239: (A) LENGTH: 2201 base pairs (B) TYPE: nucleic acid STRANDEDNESS: double (C) STRANDEDNESS: double (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 240: (C) STRANDEDNESS: do: (D) TOPOLOGY: linear алалалала далалалал алалалал ž 3 3 8 S 55 5 ន 22 8 35 8 2 2 S

1740 1380 1440 1500 1560 1620 1680 960 1020 1080 1140 1200 1360 1320 GOSCOGIUTIG CCAGCICCCA GACACIACMI GGGIAGCICA GGGGAGGAGG TGGGGGICCA GGAGGGGAT COCTOTOCAC AGGGGAGCC CCAAGGGCTC GGTGCTATIT GTAAGGAAT AAAATTITGTA GCCAGACCCC AGGIGCCTGC TCTCGTTTT CTCTGGGTGG CCTCTGATCT TETIGETETAC GTOGTGATEC CTACECGACG CTCCACGGCC GAGGCCTTCC AGATCGTCCT GETCHGGGG THIGHTGGG CACHGGGGG CGCACTTICC TGGGCACCGC CATCTTCATT GAGGCCGACC GOCGGCGGC ACAGCTGCAC GTGCAGGGCC TGCTGCACGA AGCAGGGTCC ACAGACGACC GGATTGTGGT GCCCCAGCGG GGCCGCTCCA CCCGCGTGCC CGTGGCCAAT GIGCICATOT GAGARGOTOC COCTOACOTA COTOCACATO TOCCACAGOT GGCCOTOGGG CCACCCCACG AAGGGCCTGG GCCTAACCCC TTGGCCTGGC CCAGCTTCCA GAGGGACCC ARGOGOTIGOA GODCATOAMO GGOOTIGIOTO COGGOOGITO GGOTICTOATA GIGGOGGIOÓO AGIATETEAT GTGCGGGGG ATTGCCTTCT GGTCCCTGGT GACACTGGGG TCATCCTTCA TAITITHATC TICATIOGAG AGACCTICCT GICCAIGAAC TGGGCCATCG TGGCCGACAT STOCCACCTG CTGGGTAGCCC CTACCTCATT GGCCTGATCT CTGACCGCT acaccasaac naccoccon continuo caaginocas acronacasi roncación greeneed TCCCGGARA GCATTICIGG CTGCTCCTCC TGACCCGGGG CCTGGTGGG GTCGGGGAAGG GACCAGCGGA TGACACCCGG Chaggggagg CAGATCTGAG TOTCACEGGC TCCCTGGCTC TOTGGGCTCC GGCATTCCTG CTGCGTTCCC GCGTGGTCCT nosacicate accreectea ecogaeteer ecotoregee erocitores agareacee coscioces carticoaace coossensa teceorismo tsteccams sectionism CTCTGCACCC TTCCTCTTCC TGTCCCTTGC CTGCGCCGT GGTAGCATCG TGGCCACTTA ACCGIGITICA TCTCCAGITA CATGOTOTIG GCACCIGIGI TIGGCIACCT GGGIGACAGG TACAATCGĠA GCTACATTGC GICTCATCT ACATEGABCA GITETICAAE ATEGGGGACA GIAGETETIGG GETEATECAG COGGATOCIC AGCATCTICT ACTITICACAT TCCOGTOGGC AGTOGICTOG AGGITCCAAA GIGAAGGAIA IGGCIGGAGA CIGGCACIGG GCICIGAGGG COTOGRACIOC CACTCAGATT TOCCACCCCT GAACCCCACC TOSTOGTOGG GOCICIOSCA AGAAATOCTA GITITOGICOT GICTITOCOTO GGOTITOACTO TOGGGAGACC CCACCCTGCC TTCCCGGAGA CTCCTGCTCT TCCTCTGACA TETAGGAGIG GIGGCGTTC TGCTGCTGTT CCTGGTAGTG CGGGAGCCGC CCAGTTATTC CACCATCGCG CCCACTCTCA TTGCCGACCT CTTTGTGGCC TOTOCTACAT CANTETECTS AACTACATOS ACCOCTTCAC CSTCOCTGOC TOCACCCCGT CTTCACCCCA GOGCTCCTGA AGACTGTGGG T 45 S 55 35 8 <del>6</del> S 2 15 2 22

2160 2201

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2040 2100

PCT/US98/04493

CAAGGGGAGC COTGGAGCGC CACTCAGATT TOCCACCCCT GAACCCCACC TOGTGGTGGG GACCAGCGGA SCGGATGCTIC AGCATCTTCT ACTITOCCAT TCCGGTGGGC-ÁGTGGTCTGG TEATSCITTEA TECCCOGAGA GEATTITETOG CITOCITEC TGACECOGGG CETGGTGGGG OTOCTTOCOG ACATOGAGOA GITOTTOAAC ATOGGGGACA GIAGOTOTGG GOTOATOCAG GCTACATTGC AGGCTCCAAA GTGAAGGATA TGGCTGGAGA CTGGCACTGG GCTCTGAGGG GICGOGORGO CCAGITATIC CACCAICGG CCCACICICA TIGCCGACCI CITIGIGGC TACAATCOGA AGTATETEAT GIBOGGGGG ATTGECTTET GGICCETGGT GACAETGGGG ACCOTOTICA TCTCCAOTTA CATOOTOTTG GCACCTOTOT TIGGCTACCT GGGTGACAG GTCTCATCTT TGGACTCATC ACCTGCCTGA CCGGAGTCCT GGGTGTGGGC CTGGGTGTGG OCOTOGRACIO TOGOGRAGACO COACCOTOCO TROCCOGRAGA CROCTOCTOT TOCTOTORACA CTOTOGOCCTT TOTOLOGGOC TOCCTOGOCTC TOTOGGOCTCC GOCATTCCTG CTGCGTTCCC CAGATYTOAG GOCTETGGCA AGAAATCCTA GTTTCGTCCT GTCTTCCCTG GGCTTCACTG TOACACCOOG TCTAGGAGTG GTGGCCGTTC TGCTGCTGTT CCTGGTAGTG CGGGAGCCGC TOSCCACTTA TATTITICATE TICATISSAS AGACCETECT STECATISAAC TOSSCEATES OCCIOCIDOS CICIOCACOS TICCICITAS ISICCCITOS CIGOSCOST GONAGANOS KOCCCTOGGC COACCCOACG AAGGGCCTGG GCCTAACCCC TTGGCCTGGC CCAGCTTCCA TOTOCTOTAC GIGGIGATICS CTACCCGACG CTCCACCGCC GAGGCCTTCC COGGYTECGC CACTECAACC CCCGGGCTGA TCCCCTGGTC TGTGCCACTG GTCCCACCTG CTGGGTGATG CTGGGAGCCC CTACCTCATT GGCCTGATCT GAGGECGACE GEOGGEGGE ACAGETGEAC GTGCAGGGCE TGCTGCACGA GCTCTGCGCG TTTGTTGGGG CACTGGGCGG CGCACTTTCC TGGGCACCGA GCGCCGGAAC TGGCCCCCCT CCTTCTTGTC CGAGTTCCGG GCTCTGCAGT GTOCTCATCT GAGAGGCTGC CGCTCACCTA CCTGCACATC TGCCACAGCT ACAGACGACC GGATTOTGGT GCCCCAGCGG GGCCGCTCCA CCCGCGTGCC 1020 240 180 120 300 1080 540 420 360 1440 1260 1200 960 90 840 600 80 8 5 ઝ 23 8 ᅜ 8 35 45 8 55 8 (2) INFORMATION FOR SEQ ID NO: 242: CATCHTICCC CCTCCCATCA TACCTCCTCC TTCCTGGAGC CTCTGCCGGC TTGGCTGTAA NGACAGAAAA GCAGAAGATG AGACTCTGTT CATTCACTTT TCCTAGGCCC ATCCTGTGGT CAGCCTCCTG GIGTGTATCA TAGGATTIGT TCACATAGTG TTATGCATGA TCTTCGTAAG TOGGACAGAT GOGGAGAGGA AAAAGGCAGA GATINGCCAGG AGAGGGGTGC AGGACAAAACC TOSTOSCACT TACCTOSATA TTTCAGTOSS ASSATSAAAS SCSAGACTCA CCCTACSCSS TAATTCATCA AUGITCIAGI TAATGICIAC CICAGCACCI CCICITAGCC TAATTITAGG GTTTAGCCAG CAGCTOCGGC CTCCCCGGGC CCTTGGCATC CAACTTCGCA GACAGGGTAC AGAGAGITTO GOTCAGOGGA AAAGTOTNOG GAGAAAGTOG GOTGCAGOCC CTGCAGGCCG TETTATOTEG TGATAGEACA AGTGCEAGTE GGATTGETET GTATTACAGA ATAGTGTTTT CTARGECTIT TOGITIOGGI AITAIGITIC GITTIGITAT TIGITIGITT TIGIGGETIO ATATRIATOT ATGRAARTA TAGCACIGAG GGCCCTGCTG CCCTGCTGGA CCAAGCAAAA GTTAAGAAGC CGTGGTGGTG CACCATGACA TCCAACCCGT ATATATAAAG ATAAATATAT CTOTICIAGI TECGAAGEAG TITICACIOGA AGITGIGEAG IECTOGITGE AGEITTECOG CUTOCUTUCT GUTOCUCCUT CUCUAGUCUA CITISCUUBAG TIGIGUTIGU CGCIUCITAI TICICICITI CICICCICCO CACCICICAC COTIGOCOTO ICCAICICO ICICCOCOC AGGITGCCCA ATTITOTITC TICAATTITA CIGGITACIT ITTIGIACAA AICAAICICT ICCCCICCIC CCICIOGCIC CCCGICICAT IICIGICCAC ICCAIICICI CICCCICICI ANCIOCCTIC GITTICGIOTA GAITGACGCG TITICITITGIA ATTICAGIGI TICIGACAAG E Ě SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 242: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear € TYPE: nucleic acid LENGTH: 1146 base pairs

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480 420

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240

180 120

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AGATCAGCCG

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TYPE: nucleic acid TOPOLOGY: linear STRANDEDNESS: double LENGTH: 1661 base pairs

SEQUENCE DESCRIPTION: SEQ ID NO: 241:

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INFORMATION FOR SEQ ID NO: 241: (1) SEQUENCE CHARACTERISTICS:

WO 98/39448

1080 1020 840

960 900 720 660 600

780

CCTCTGATCT TOCACCCCGF CTTCACCCCA GGGCTCCTGA A

GIAACOGAAT AAAATITOTA OCCAGACCCC AGGIOCCIOC ICICGICTIT CICIOGGIOG TOGGGGTCCA GGAGGGGAT CCCTCTCCAC AGGGGNCACC CCAAGGGCTC GGIGCTATTT 1661 1620 1560

GAGGGACCCT GGGCCGTGTG CCAGCTCCCA GACACTACMT GGGTAGCTCA GGGGAGGAGG

1500

470

WO 98/39448

1320 1350 0801 1140 700 1260 960 020 22 8 240 8 90 20 480 540 8 8 22 780 840 90 1146 GCTGTGCCAC GGCTGTTGCT TCGGTTATTT AAATAAAAG AAAGTGGAAC TCGAAAAAA AGTECECCCA TCAGGCCACA CTGCTGCCAC CTCTCACACG CCCCAACCCA GCTTCCCTCT TETETGGGAE TTGGAGAGAE ATEACTAACT GATGGCTCCT COGTAGTGCT CCCAATICETA TOSCCATGAC TOCTGAACCT GACAGGGGTG TGGGGAGTTC ACTGTGACCT TECTECOSECA GECATAGAG GECAGAGESC CECTGAGAGEC CAGTTGCAAA CANGGETCAG AGCAGAGGG GEGGAAACCC GTTCGCCGAG CCCAGCGAGC TTGACAACCC TECETECITIG CAGCECTEGA GAAAGETEAG CECCACAGAA CETAAGAACT ATGGETEATA CONCONTING ACTOCORGO CONTINUISMO CTGGTACCGC CCCATGTATA AGGCTTTCCG ACCATGTGCT CITHERCOTC CAGGCCATTG GIATCCCAGG TTGGGGATTC AGTGGCTGGA TCTCTGCTCT SCHOOLGCOG AAGGCAACAC AGCAGTATCC GTGCTCATGC TGCTGGTCGC CCTGCTCTTC ACTECCATTG CTGTGCTAGG AATTGTCATG CTGAAACGGA TCCACTCCTT ATACCGCCGC ACAGGIGCCA GCITITCAGAA GGCCCAGCAA GAAITITGCIG CIGGIGICIT CICCAAGCCI GEOGTICEGAA CEGEARETTG CEAATGEAGE CGETGGGGET GETGAAAATG EETTEEGGG COCOTGACCO CTGACTOGGA TOCCOTGGCC CTGCTACTTG AGGGACCTGA CTTAGCTCCC CITICAGCCA CCACCAGCCT ATGAGCCTCC AGCCCTGCC CCATTGCCTC CACCCTCAGC crececreses ROSCACAGOT ACTOGACAGA ACAATTOGOC COCTOTACOT TOTTTTTOTO CAGTICAGOS CTGTATCCAC CANGIACTAC CICTIGGATGT GCAGCAGGST GGCTCTTCTC CIGAACTICC TGGCCTGCCT TOCTICHOGGI GAGTGACAGT TCATTCAATT TCTTCGTTTT CITCTTCATT TTCTTCGTCC CAACCOGAAG GCAGAGGAGT TOGACCGAAG GAGNCGAGAG CTGCAGCATG CTCCTTTTTC CAGGACATCT CCATGGAGAT CCCCCAAGAA TTTCAGAAGA GOCCAGCTTC TGTGTGGAAA CCAACAATGG CGCAGGCTTT GGCCTTTCTA SEQUENCE DESCRIPTION: SEQ ID NO: 243: i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1150 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear 69 (2) INFORMATION FOR SEQ ID NO: 243: AAAAAAAAA AAAAAAAAA GGGGGNCCNC ź STCCCTANGG 3 **AACCCACGGC** CARTIC

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INFORMATION FOR SEQ ID NO: 244: (5) S SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1529 base pairs

TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear 3 E 0 6

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SEQUENCE DESCRIPTION: SEQ ID NO: 244: ž.

8 120 180 240

300

360 420

TOCTAGGITT CATTITICCCA TITICSCAAAT CACCTICGAA GICTICGCCCC ATGACAGCAA AGRANGSCY TANGGCGGCC CACTCACTAY GGYGGGCTGA GTGGAAGGTC CYNAACCAYG TCCCAGAGGC COGGGGGTTC CAGCTCTGCC TGTAGCAGAG CCCTGAGGAG GAGGAAG AGAIGICI GAAATACGIC CGGGAGAICI TITICAGCIA GGGCATAAAC IGIGCACIGA **1666000MCC** OCCIONOSCA ANOSOCIET CIGNOCIEET ACCITITATOC CITETIAGGE CIGGEAGATI CACCTCAGGC CAGAAGCCCC TGGACACTCC GGGCCTTGGG GTGCCGTTCT GAGTGTGCGG AAGGCAGGAC TCAAAATGAG ATCCCATTTG ACTCCCTCTG TATGTACTGT GCCCTCTC TTCCAGGGAA TGCAAAGAAA TCTCCTGAAG CCCACCTCTC CCACCTTCAC GAGCCCAGAA THICAGIGIG GCTTTGGACA GGAATATATG AATAAATCAC TGCCATACAG GTTTTCCAAT ACACAGTGC AAGTTCTGGG GIGCGCAACC GICCAGICCT GITCACAGGG CHETCHOSOT TCAGGTCCAG GGAAGCTITIG AAGCAGTCAA GCCTTGTCTT TGTACCCCAI TACCTIGCT TCCCCAGAGA CACTGAGGTG CTCGCTCTTT TAATGTCCTC GTTTGTTGCC GTAAGTTCT GIGIECCIGIC TITICITICAGI CACTCAGAGA TCACTCCTGG ACCTCTGGGG TITGGAGTIT TAGAAATAC ACACAATTCC CCAATGCGTA AGTTGTGCTA ATGTCTTTTCC ACTIGITCTICC GAGAGGACAGC TGAGCTTTCCA CTGGTGCTGC OSCIPCINGAG SCIPCINGAGI COCAATION TONGITAGIC AGNOACCAGG AATRATGTCA TGTGGTGGTC CAACTTACTG GAACCAAAGA GACAGTACTT CCGCATGATC GAGGAAGCCT GTTTCAAAAA TAGITTCCAT CATGAGTCTA TCAATGAGCT CAGCCAGCCT AGAAAGCAAA CGAGCTGCCC ACAGTTCTCT GCCCTGTCTG GCCACAGTOT ATAGACTOGT AAGCCAGACA GGCCTCCTCC CGCAAGCTGC TCACCTGTAC CTTGGTCCCC GGGCAGCTAG CTATAAAGCA AGAGGGACAG GGGAGTGATT TGTCAGGGGA GAGACACTGA GGACAAGAGA TCACACCAGA GTACATGTCT CTGCCTCTGT AGGATCACTG CCAGGTGCAC TGGAATTGCT ACAGTTTAGT AGGATTTGGA GICCTCCAGG GICTCATCAT ITCGGAAGIG GAGGGIGGCA GCGITITGITIT 45 တ္တ 55 15 റ്റ 25 8 35 **4** 

720 780 840

900 99

480 540 1140 1200 1260 1320 1380

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8 SS 8 дисандате ессеснасе сеснасавсе тессавсее вазовате снанасими CTGAAGGACA AACAATCTTG TACTAAGAA CAGGAATTGG TCACTCCCTC CCCACCCTCC TGAAGCTAGA GGAAGATTTG CTCAGATCCA TTAATTAAAG 2 CCACCTICAC ACTITICANA CIGANATICA ACACTITICA GAGACCCIGA AICCITICICI GAGGCCCCTG AGAGCTCCAC CTAGTTCACA GGATAAAATC CCACAGCAGA ACTCGGAGTC AGCANTOSCT AAGCCCCAGG TOGTTGTAGC TCCTGTATTA ATGTCTAAGC TGTCTGTGAA ACTOTYTOTO GGAGAACTTY AUCTYAACOT GGAGATCAAG GGAACAAAIG GACAGGTIAC TAGCCCACAG AGTGGCAACT TCCGCCAATT GCTACTTCAA AGATGTCGGA CTGAATATGA CCCAAATTTC TCTTATATGG GAGCTCGCCT GTGTAATTAC CTGTCCCATC ATCTGACAAT TACAACAGAT GATGCTTTGC AAGAACTTGT GGAACTCATC TATCAACAGG CCACATCTAT TIGRCAGGAT TATCCTACIC TATCAGAATA TGTTCAGGAT TYTTTGAARC ATCTTACAGA TOCCCCTGAA TITTACCCTT CAGGITAITC TICCAGITAC ACAGAAICCT AIGAGGAIGG ACTTANAGAT CAACCTGCAA AAGGGGATGA ACTTACTCGA AAACGATTTC ATGCATTTGT TATOGATGAC AATTIAATTT GIGCAGTAAA ATTOTTAAAG TIGACAGGAT CAGITTIGGA AAGAGCAGAT ATTCTTCAGG TIGGICTICG AGAATIGCIG AAIGCCCIGI TIICTAAICC TITCACTUCA GCTGATCCAG ATTACCAAGA GAATACCAA GAATTACTTG AAAGAGAGGA AGAMANIGAT CCTAACTACT TINIGANIGA ACCAACNITI INTACNICIG AUGGISTICC ACTECCOGTEA AGTAACTGGG GCAGAGTECA TOCAACTTCA ACATATAGAG AAGCAACACE COTTOTCCTA GATOCAAACT GCAGTAGAGA TOTAAAACAG ATOCTCTTGA AGCTTOTAGA AGATOCTTOG AAGGAAAAAG GAAAGATGGA TATGGAAGAA ATTATTCAGA GAATTGAAAA GGANTCAGAG COTAAGCGAA AACAGTAAAG TTAAATTTCA GCATATCAGT TTTATAAAGC GCATCATATT GATGATGAGA TGGACCCAGA GATAGAAGAA GCTTATGAAA AGTTTTGTTT CTITITITCCA GATTATGAAG AAAATGGAAC AGATTTATCC GGGGCTGGTG ATCCATACTT INFORMATION FOR SEQ ID NO: 245: Ξ TOTCACAATO AGCTOCATOG TITTAGGGAGT CTTTGGGAGC CTTGGAAGTC SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: (B) TYPE: nucleic acid (A) LENGTH: 1537 base pairs STRANDEDNESS: double TOPOLOGY: linear 1500 1440 1529 540 360 300 240 180 120 1020 480 420 1080 660 600 6 1260 1140 90 840 780 720 960

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GCCCCTOTGA GOOCCAGCTC TOGAAAAACC TOGGAGTTGA TOCCOGAGGY TOGGAAGAAC ATUACCCCTT GUAACTGTGC CGAGTTCCTT AAATCTCAGC TGGGATCCTG GACCTGGGAG GICTITCITT INCIGITITG AGTIGGIGAG IGAGIGAATA GGGIAACAIG GGCCITCAGG

180 120

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SEQUENCE DESCRIPTION: SEQ ID NO: 247:

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3 23 20 ᅜ 5 ઝ 25 8 50 CARATTRAGG AUGITGAGIT AUGITACIAA TOTAUGCAAC TITAATTITUG TITAACACIA TCTGCCAAAA TAAACTTTAT TCCCTATAAC TTAAAATGTG TATATATATA AGTITIAGGTA TOGTGATITTA GCAGAACACA AGAGAGCAAG AAAATOTGTC ACATCTATAC AMAMAMAMA AMGOGNOGCC OCTOTAGAGG ANCOMAG TTATISTACAG TTAATICTAC ISTITISSCT SCAATAAAAT CSATITISAA ATAAAWRAAA TOCHGANTIT GOCCHGANC CSCCGCGGTG GCGGTTGCTA TCGCTTCGCA GHACCTACTC (2) INFORMATION FOR SEQ ID NO: 246: ATATTACTTT TINGTITGAT ACTANGIATT AAACATATIT CIGKATIATI CCAAAAAAAA ATCCCAGCGG TCCTTACCAG AAAAAGCCTG TGCATGAAAA AAAAGAAGTT TTGTAATTTT TIGICACAGE AGIATGETOT CITICEGACG GGGCCCTIAT TIACCGGAAG CITETGITICA CTOTOTTOGC ACTGATACCA GAAACCACAA CATTGACAGT TOGTOGAGGG GTGTTTGCAC AGATOCTOCO OCTOGATATT ATCAACTCAC TOOTAACAAC AGTATTCATO CTCATCOTAT ATRACOTOCA OCCOARAATA AAACATCOCC CCTTCTOCTT CAGTOTGAAA GOCCACOTGA ACCEMBECAG ETGAGAAGAG TIGAGGGAAAA GIGEIGETGE IGGGIETIGEA GAEGEGANIGG aaaaaaaaa aaaaaaaatt togtog (2) INFORMATION FOR SEQ ID NO: 247: E X Ξ (A) LENOTH: 506 base pairs
(B) TYPE: nucleic acid
(C) STRANDELNESS: double
(D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 246: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear LENGTH: 1348 base pairs TAATAGTTTA 1380 1537 1500 1440 1320 180 240 120 360 300 480 420 908 8

WO 98/39448

	TCTOCTICANG GOCAGOGTISC CCTGGAACAC TGGTAGTTCT GOGGCTGGGA GGGAGAGGG	240
٧.	CTCCGGCTIT CTCTGAAATG AACACTGCTC TTCAGCAGTT CAAGTACTTG TTCTCAAAAC	300
<b>,</b>	ATTITCTAAT TGATTGGTAG GTTTTCATAA GCATTGTTTC TTTAAGGCAT GGAAAGGGAA	360
	GAATGCTCAA GCAAGTCATG TITGTTTTCA GTGGGATGGG CCGGGGTTCT CACTGCTGGG	420
01	GOCTICCCCT TOCANGIBGC ACCTITISTICC AGGOCCACCA GOCAGACTCT TCCCACCTTC	480
	TCCCACTGAA GCACCAAGGG GCTTGAACCG TAAITTTGGCT AATCAGAGGC ATTTTTTTG	540
7	TCCTAGTATC TITCACACTT GTCCAACCGT CTTATTTTTT TAAAAGTTCT GTTGCTTGTA	009
<u>,</u>	TTAACAGGAA ACTAGAGAGA AATAGTTTCT GAAGCCAGTT TATTGTGAAG ATCCCCAAGG	099
	GGAGGTTCGG TAGAGAAAA TAGTAAGCTG GTTTAGAAAC TGAGGAGGGC AAACAGCCAG	720
20	GACGCATTGG AGAGGAITTT GCCAAAGAIC TACCCTGAGA TAAGGCCTGT CCAGTGTCTT	780
	CACCACOTGA ATAACCAGGG CTCCAAAGTG TTTTTCTGCT TTGAAAAAA AAATTCCACA	840
2,5	AGCTTTTAAA GGTGCATTTA AGAATCCATG TGACTTTAGA ATGGAACTGC CGGCCCTGGC	006
3	AACTOTCAGG TOTOCTAGAA GOTTCGATGC CTCTGGAATG CATGTGATAC TCATCTCCAT	096
	TITISTITICS TANTISCAIT TITISTICITY TAGCAGATCT STCCCTGTGG STGGTGTCTA	1020
30	AGAAGTOGGA CACCTTOGTT TITGTGTTAG ATTGAGGTOG GCAGCTGCAA TCAGCTTCTT	1080
	TATATOCAAA TTAGGCACCA CCCATCTGTG GTTCCCTGGT TGGTGGCTAA TGAAGTGAGG	1140
25	GGAGGGAGGG ATGTCACCCC AAAAGTAGGC CCTCCCATTG GCTTTGGCCA GGCCAGACAC	1200
3	TICACATCOT TTACATGGTT CTGTGTAATT TTAAAGTTTA TGTGTATAAA GCGAAGGTGT	1260
	tictgigada cigtatattt tgtaartaaa tatattgcta cittgagawr aaaaaaaaa	1320
40	AAAAACTCGA GGGGGCCCG GTACCCAA	1348

INFORMATION FOR SEQ ID NO: 248: 3 45

(A) LENGTH: 1766 base pairs (B) TYPE: mucleic ecid (C) STRANDEDMESS: double (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 248:

8 120 180 OTGCCGAATC GCCAGAGCGG CACGAGCGGC CACGAGAGCA GGCGAGTAA AGGGACTTGA CACCETTETE CCACCETOGC TEXOSTASCA TEXESCAGOC TEXECOSCEA CTEAGTECEA GOGAGOCAGY TOCOGGATTA TYCTATTYCC CCTCCCTCTC TCCCGCCCCG TATCTCTTTT 8 55

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300 360 420 480 540 9 9 720 780 840 900 960

1500 1020 1140 1440 1560 INCLAICTOC TOSTCOTOCT TOGAGOCGA GCOSTCOGCG CCCGGCGCG GCGGAGCCC CICTIGICACC ATCACCITICA GGATCIACAA GICCGICATIC CAAGCIGIAC AGAAGICAGA CCATAATTAC ATGAATGCTG CCATGGTGCA CATCAACAGG GCCCTGAAAC TCATTATTCG AGAGCOTICO COCGCOCTIGG GCACGAAGAG CTGCAGGTICO TOCTGTGCGG, TGCAGGATO CATTITICIOS AGAGATOTOS AGAAGACTOS GITITOTOTIT GOCACCACOC IGAICAIGO AGAAGGCCAT CCATTCAAAG CCTACCTGGA CGTAGACATT ACTCTGTCCT CAGAAGCTT GACCTATGIT GGICCIGITI TIAACGGAAT CACCCITICIA AITCITGCIG AACTGCICA TITCAGIGIC CCGATIOICT AIGAGAAGIA CAAGACCCAG AITGAICACT AIGIIGGCA CAAAAAAAG GCAGAATAAG TACATGGAAA CCAGAAATGC AACAGTTACT AAAACACCAT TCCAGTITIC AGCACTAGIC TIACTCAGCT ATCCAITATA GITTIGCCCT TAAGAAGIC TIGGAGCCCI CAANTCCIAT CITCCIGCCC CACAAIGIGA GCAGCIACCC CIGAIACIC CTGATTCCAA GAATGCCATC TGATAAAAA GAATAGAAAT GGAAAGTGGG ACTGAGAGG AGTCAGCAGG CATGCTGCGG TGGCGGTCAC TCCCTCTGCC ACTATCCCCA GGGAAGGAÀ GOTTICCOTG GCAGOTTICA GIGICATCAG TGIGGITTICT TACCTCATCC TGGOTCTIC TCTCTTTCTG GTAGAAGAIC TGGTTGACTC CTTGAAGCTG GCTGTCTTCA TGTGGCTGA COCCCOAGAT CAGACCAAGT CAATTGTTGA AAAGATCCAA GCAAAACTCC CTGGAATCG TIAATAGTIA TAACGICGIT ACTIGIACIA IGAAGGAAAA TACICAGIGI CAGCITIGAG AAGAATCAAA TTCATAGGAT AAGTCAATAC CTTAATGGTG GTAGAGCCTT TACCTGTAG TICADAGGG AAGGATIGGA GGIAAGAGAG AAAATGAAAG AACACCICTG GGICCTICT TITICITIAA TGAITIAACI AICAACIIGA TAAAIAACIT AIAGGIGATA GIGAIAATI CIGCATICCA AGCITITITI TIAATITISSI GITTICICCC AICCITICCC ITTAACCCI AGIATCAAGC ACAAAATTG ATGGACTGAT AAAAGAACTA TCTTAGAACT CAGAAGAAG TGATTAACTT ATGAAAAAT TATTTGGGGA CAGGAGTGTG ATACCTTCCT TGGTTTTT ဓ္က 35 8 45 2 2 2 23

1080

1200 1260 1320 1380

CCCAAATCGC CGGATATGAT CGTAAA

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1766

1620 1680 1740

RECTCCSCCA TITIOGGAAAG TGGTTTCTAC GTCACTGGAC ACCGGTTCTG AGCATTAGTT TORGARCTEG TICCEGRATE TECTITECTE CETETECET GECCACETEA AGITTAATAA ataaggitigt acttittctta ctataaata aaaaaaaa aactcgaggg gggcccggta

(2) INFORMATION FOR SEQ ID NO: 249:

(i) SEQUENCE CHARACTERISTICS:

9098 STRANDEDNESS: double TYPE: nucleic acid LENGTH: 2664 base pairs

TOPOLOGY: linear

S AGTOTOCTOG GAGCAGGOGG AGTAAAGGGA CTTGAGOGAG CCAGTTGCCG GATTATTOTA (xi) SEQUENCE DESCRIPTION: SEQ ID NO:

5 20 2 25 ассамассот ссасасссва савсавсава массемама сетоссскае сетоваамсь TASCATORCE GARCETCORC GROCACTICAG TOCCATTOCA TOTOCTOCTIC GIOCTTORGA THICCCCICC CICICICCC CCCGTATCT CTITTCACCC TICICCCACC CICGCICGCG AAGAGCTGCA GCTCCTCCTG GACTOCTTGA AGCTGGCTGT CTTCATGTGG CTGATGACCT ATGTTGGTGC TGTTTTTAAC ATCAGTGTGG ACTROOTITIG TETTTGGCAC CACGETGATC ATGETGETTT CECTGGCAGE TITTCAGTGTC TACAAGTCCG TCATCCAAGC TITCTIACCT CATCCIGGCT CTICICICIG TCACCATCAG CTICAGGATC ACAGGGCCCT GAAACTCATT ATTCGTCTCT TICTGGTAGA AGATCTGGTT ACATTACTOT GTCCTCAGAA GCTTTCCATA ATTACATGAA TGCTGCCATG TGTACAGAAG TCAGAAGAAG GCCATCCATT CAAAGCCTAC TOCCOTICCAC GATCTGATTT TCTCGAGAGA TCTGAAGAAG 600 540 480 420 360 300 240 180 120 660 60

35 ઝ GGAATCACCC TICTAATTCT TOCTGAACTO CICATTITCA GIGTCCCGAT TGTCTATGAG CTTCHARAGA TCCARGCARA ACTCCCTGGA ATCGCCRARA ARARGGCAGA ATRAGTACAT AAGTACAAGA COCAGATTGA TCACTATGTT GOCATCGCCC GAGATCAGAC CAAGTCAATT GGRARCCAGA AATGCAACAG TTACTAANAC ACCATTTAAT AGTTATAACG TCGTTACTTG 900 840 780 720

TACTATGAAG GAAAATACTC AGTGTCAGCT TGAGCCTGCA TTCCAAGCTT TTTTTTAAT 1020 960

CTGATAAAAG AACTATCTTA GAACTCAGAA GAAGAAAGAA TCAAATTCAT AGGATAAGTC

CAGCTATICCA TTATACTITT GCCCTTAAGA AGICATGATT AACTTATGAA AAAATTATTT 1320 1260 1200

GOGGACAGGA GIGIGATACC TICCTIGGTI TITITITIGCA GCCCICAAAT CCIATCTICC TOCCCCACAA TOTGAGCAGC TACCCCTGAT ACTCCTTTTC TTTAATGATT TAACTATCAA

CTTGATAAAT AACTTATAGG TGATAGTGAT AATTCCTGAT TCCAAGAATG CCATCTGATA

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TUTACOTCAC TOGACACCOG TICTGAGCAT TAGITTGAGA ACTOGITOCC GAATGIGCTI GTEACTECET CTGCCACTAT CCCCAGGGAA GGAAARGCTC CGCCATTTGG AAAAAGAATA GAAATGGAAA GTGGGACTGA GAGGGAGTCA GCAGGCATGC GAAAGTGGTT

1140 1080

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SEQUENCE DESCRIPTION: SEQ ID NO: 250:

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(2) INFORMATION FOR SEQ ID NO: 250:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 865 base pairs a) TYPE: nucleic acidb) STRANDEDNESS: doublec) TOPOLOGY: linear

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AAAGGTCGTG ANTGGGGGAA ANCC

COTOCATAGO TOTTTOTATA GOGGICCOCC AMATTOCATT CANOGGGCCG TOGGITTITAN

2640 2580 2520 2460

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actigiigita cototictigiig cotocaacat aaaaatacag tagcacctaa ggagctitgaa TOCCACACAG GATTITITIT TITITITAAGA AAAACCTATA GATGAAAAAT TACTAATGAA TGCCACATCC AGTICTITTC TITTGTTGCT GCTGTGTTTA GATAATTGAA GAGATCTTTG CCAGGAAGIC AGIGATIOIC TITTIOGOCT TCCCCTCCAA AGGACCITCT GCAGIOGAAG

2400

2340 2280

aaaaaaaaaa aaaagggcgg ccgctctaga ggatccaagc ttacgtacgc gtgcatgcga

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GTITTIAACTG GTTTCATGTC CTAGTAGGAA GTGCATTCTC CATCCTCATC CTCTGCCCTC

2220 2160

TIGOATOGGT AAAAGGIACC CTIGCCTTAC TCCATCTIAT TITICTIAGCC CCCTTIGAGI

CAGCTOCAAA AAGTAGTOGA AGGAAATTOT CTACOTOTCT TOGAAAAATT AGTTAGGAAT GGAATATGCA CTOOCGAGIT TAAAGTAAGG GCTATGATAT TIGATGOTCC CAAAGTACGG GAAAGAAAAA TCATCTOTGA OTTCCTTCAG OTTCTCACTC ATAOTCATGA TCCTTCAGAG

2100

2040

1980 1920

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CGATTCCCAT TIGGGGGCAA GITTTITICT TCACCTICAA TAIGAGAATT CAGCGAACIT

CATGIGGGCT CCTCAGITAT TGAGITITIG TGAICCTAIC TCAGICIGGG GGGGAACATI

AMATAMATOT CTOTAMCTOC TOTOCACTOC TOTAMACTTG TTAGAGAMAA AMATAMCCTG TODICCION COCCIOCOCA CONCAAGITI AATAAATAAG GIIGIACITI TOTTACIATA

1680

CICAMBMOST GAMMIACAGA MAGCCITITI TICITGATOT TITICCCGAGA TICAMATOTO

1860 1800 1740

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8 . 45 THOGRETTET CHECCARECT TICCETTIAA CECTEAGIAT CAAGCACAAA AATTGATGGA AATACCTTAA TOOTOOTAGA GCCTTTACCT GTAOCTTGAA AOOOGAAAGA TTOGAGGTAA

GAGAGAAAAT GAAAGAACAC CTCTGGGTCC TTCTGTCCAG TTTTCAGCAC TAGTCTTACT

1380

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TIGOCGITIC CICOGOCTY IGGIGGGAIC GGIGICCICA GGAIGAGAIT IAGGGITICC GOOTICCTICCO GATTICAGGTC CCCGTTCCTA ACGOTGGGAT CGGTGTCCTC GOGATGAGAT COTOGGAGTO AGOTACCAGA TICAGCCCAT TIGGCCCCGA CGCCTCTKTT CICGGAATCC

TOGGGGCTTT CGGGATCTTC ACCTAATATC CGGACTGCAA GATGGAGGAA GGCGGGAACC

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TAGGAGGCCT GATTAARANG GICCANCTAC TOGICTIGIC AGGIGCCIGG GOCANGCAAA

300 240 180 120 69 PCT/US98/04493

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	TOTGOSTGAC CITICOTCICA GACITICCTGC ITTTCCGAAG CCITICCCCGA CAIACCITICG	360	CERTAINING MAINTAINE OF THE PROPERTY OF THE SERVICE
٠	GACTAGTGCA GAGCAAACTC ITCCCCTTCT ACTTCCACAT CTCCATGGGC TGTGCCTTCA	420	CONTRACTOR OF THE CONTRACTOR O
8	TCAACCTCTG CATCTTGGCT TCACAGCATG CTTGGGCTCA GCTCACATTC TGGGAGGCCA	480	S ACTIVITIES CHARACTER CARACTERS CONTROLLED TO ACTIVITIES OF THE CARACTERS
	GCCAGCTTTA CCTGCTGTTC CTGAGCCTTA CGCTGGCCAC TGTCAAGGCC CGCTGGCTGG	240	ACTIONAL GENERAL CARLINAMES CALLESTING CAMMENTAL CHAMMENTAL CALLESTING
9	AACCCCGCAC CACAGCTGCC ATGTGGGCCC TGCAAACCGT GGAGAAGGAG CGAGGCCTTGG	. 009	ACCIOGRAFIA CONTRATA TRICTITITA ATAGAMICA COCCOCACTI  10 TOSCOACTA AMPRETICADA TRICTICADA COMPRIMENTE TABABAGICA
₹.	OTGGGGAGGT ACCAGGCAGC CACCAGGGTC CCGATCCCTA CCGCCAGCTG CGAGAGAAGG	. 099	
	ACCCLANGTA CAGTOCTCT COCCAGAATT TCTTCCGCTA CCATGGGCTG TCCTCTTTT	720	THEOREMS INVESTIGATION INTO THE PROPERTY OF TH
15	GCAATCTIGGG CTGGGTCCTG AGCAATGGGC TCTGTCTCGC TGGCCTTGCC CTGGAAATAA	780	15 metablica remaining manager and manager man
	GGAGCCTCTA GCATGGGCCC TGCATGCTAA TAAATGCTTC TTCAGAAAAA AAAAAAAAA	940	COLONGO DE MANAGERA DE LA COLONGO DE LA COLO
70	AAACTOGAAGG GGGGCCCCGGT ACCCA	965	20 TGATGTTAAG GGGGGTAGAG ITTGCAAGGG GACTGTTTAA AAAAAGTAGC ITATACAAGG
			TOTOCTTOCA ACTTABATAT AGGTOGGTA TOTOTAGTOT TISCTATACC ACTGACTGTA
č	(2) INFORMATION FOR SEQ ID NO: 251:		TICAAAACCA AAGINITIAAG AQQQQAAACG CCCCTOTITIA TAICTGTAGG GGINITITIAC
3	(1) SEQUENCE CHARACTERISTICS:		ATTCAAAAAT GTATGTTTTT TTTTCTTTTC AAALTAAAG TATTTGGGAC TGAATTGCAC
	(A) LENGTH: 2082 base pairs (B) TYPE: nucleic acid		TAAGATATAA COTGCAAGCA TATAATAGAA AAAAAATTG CAAAACTGTT TAGAAGGCTA
30	(C) STRANDEDNESS: double (D) TOPQLOSY: linear		30 ATAAAATITA TGCAGITATA AAAATGGCAT TACTGCACAG TITIAAGATG ATGCAGATİT
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 251:		TITTACAGIT GIAITOTOGI GCAGAACTOS AITITCTGIA ACTIVAAAAA AAAICCACAG
35	TGGGGGGGN ANTGGGTGTC TGGCTCANGG ATTGCCNAAT CTGGAAATTC TCCATAACTT	09	TITIAAAGGC AATAATCAGT AAATGTTATT TICAGGGACT GACATCCTGT CTTTAAAAAG
3	GCTAGCTIGT TITITITIT TITITITACA CCCCCCCCCC CCACCCCCCG ACTICCACAA	120	AAATGAAAAG TAAATCTTAC CACAATAAAT ATAAAAAAT CTTGTCAGTT ACTTTTCTTT
	TOTTCAATGA TOTCAGCAGA GITOTTCATG TGAAAGGTTG ATCACCTTTG AAGCOTGCAT	180	TACATAITIT GCTGTGCAAA AITGTTTTAT AICTTGAGTT ACTAACTAAC CACGCGTGTT
40	CATTCACATA TITITICITC TICTTCCCCT TCACITICATG AACTGOTGIT CATTITICIGT	240	40 GITCCIATGT GCITTICTIT CATITICAAT TCTGGITATA TCAAGAAAAG AATAATCTAC
	STOTOTOTOT STITIALITY STITIGARITY THIPPITAA THINACTITY AGACCITISCT	300	aataataaac ggcatttttt tttgaaaaaa aaaaaaaaa aa
4	GIOTIGOCCA CCTITITICC AACCICCACC CICACTCCIT CICAACCCAT CICITCCAAG	360	
}	ATGAMGAAA AAAAAAGCA AAGTITTITI ITCITCTCCT GAGTICTICA IGTGAAATTG	420	(2) TAENDRAMMENT FOR MY AND ACT.
	ACCTICCAAA GGAAAAAAA AIGIGAAAIG ITAIAGACIT GCAGCGICCC GAGITICCAIC	480	. : 10 TOWN TOWN TOWN TOWN TO THE TOWN TOWN TOWN TOWN TOWN TOWN TOWN TOWN
20	GOOTTITIT TITIAGCATTG TTATGCTAAA ATAGAGAAAA AAATGCTCAT GAACCTTCCA	540	50 LINGTH LENGTH AS Deser pairs
	CAATCAAGCC TOCATCAACC TTCTGGGTGT GACTTGTGAG TTTTGGCCTT GTGATGCCAA	009	(c) 17PE: INCLETE ACLO (C) TOWNDENESS: double (f) TOWN OCY 11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
۲,	ATCHGAGAGT TTAGTCTGCC ATTAAAAAA CTCATTCTCA TCTCATGCAT TATTATGCTT	099	
3	GCTACTITIOT CTTAGCAACA ATGAACTATA ACTOTITICAA AGACTITIATG GAAAAGAGAC	720	ביינים מיינים
	ATTATATTAA TAAAAAAAA AAGCCTGCAT GCTGGACATG TANGGTANAA TTATTTTTC	780	בייסיפיים מכנינימסס שהזדרונורו מפנינומיור הרזנימשפתה בזבתפכתפונו.
9	CITITITITI CCTITITGCT TGGAATGGA CGTTGGAAGA CTTATAGCAT GGCAITCATA	940	GUCUGCUTGG CCCCTAGGGA GTCCTTAGGC AGGATGGAGG CTGTTGTGAA CTTGTAGCAA

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45 6 ઝ 30 25 20 2 5 S GGAAACCAAA AAAAAAAAAAA AAAAACTCGA GGGGGGCCCG TA TOCCCTCATT GCTCTACAGG TICTOGTATO CCAGTCATTA TICACCICAT CIGGATGTAT GGCACCATCT ICTICATGCT GICCIGGICI GIGGCACAAC CCTACCTIIG GIGGAAAAAG CACAIGACAG CCAITCAGCI GAICCAGITT TODACCIOCI CTCCGAAAGA AAGACGGGCA GOTGACCTTC CTACATOTCT TCCATCACTC TGTGCTTCCC GIGGCACTOT COCTOTACAT IGICIAIGAG ITCOTGATOT CGGGCIGGOT GAGCACCTAT AUGGENANTE GERAGESETT SCAGETISSET GESTTSCATER TIGISTREAR STISTERSET CTAATGACCT CCATTCTCCT GACCTACGTG TACTTCGTTC TCTCACTTGG GCCTCGCATC CTOGGAATAC AGCCTGTGGA GGCTGCTTAC TCAACTTGTG TCTTAATTAA AAGTGACAGA CCCCTCCCTG CCTTAAAACT TOGGAGAGGA GCACTCAGGG CTGGCCCCAC AAAGGGTCTC ATAMACICTT TOCTOCACOO CACACACTGA AGCAGTAGOT TOTGGGCCAA AGGTCAGGGT GGGCGGGGGC CACTGCACAT CTCCCAGTAC TACTTTATGT CCAGCTGTAA CTACCAGTAC TCCTCACACA GAAGAGGTCA GCAATAATOT CACTGTGGAC CCAGTCTCAC GCTGTCTGCC ACTCCAGAGC TGGGGGCTAA AAGGGCTGTA CAGTTATTTC GACCACGGCT TIGGTICCTC ACCCACTICC CCCGGGCAGC TCCAGGGATG CITATOTOGI CAGGACTGAG CAGGGGACTG GCCCTCCCCT CCCCACAGCT AGGACTICAC CITAGGGCAG TOTOCOTICAG TOCCOTOTICO ACCTACACOT TROCCAAGGT ACTOTTATAC CAAGGGCAAG CGGCTGCCCC GTGCACTTCA COGRECATOR CATAMIGIAC CIGRACIACE GAIRATCIEC CITIEGECCT TOTTCCTCTT CTCCAAGTTC ATTGAGCTGA TGGACACAGT GATCTTTATT GTGACCCTGT GGACTATTCC AACAGCCCTG AGGCACTTAG GATGGTTCGG CAAGGCCAAC TGAGAAGCAT GGCCTAGATA GGCGCCCACC GCAMATIGGA GITTCTCCAAC 1260 1200 1140 1080 1482 1440 1380 1320 1020 960 900 840 540

780 720 660 600

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GAGGTGATGA AGCACGCAGA TCCCCCGGATC CAGGGCTACC CTCTGATGGG GTCCCCCTTG

480 420 360 300 240 180

2 INFORMATION FOR SEQ ID NO: 253:

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Ξ SEQUENCE CHARACTERISTICS: 9 O B € TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear LENGTH: 834 base pairs

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ž SEQUENCE DESCRIPTION: SEQ ID NO: 253

8 GGCACGAGCG CCGTTGCQCG CCTGGCCCCT ACGGAGTCCT TAGCCAGGAT GGAGGCTGTT

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TOOCTUAGEA CETATACETO OCOCTOTGAS SCITCAGGACT GEASCETTAGG GEAGTOTECO CCAMAGGICA GOGIGGGCGG GOOCCIGGGA AIMCAGCCIG TGGAGGCIGC TIACICAACI ADDOCTOGEC CCACAAADOG TETEGTOGEC TTTTTTCCTCA CACAGAAGAG GTCAGCAATA CTAAAAGGGC TGTACAGTTA TTTCCCCCCTC CCTGCCTTAA AACTTGGGAG AGGAGCACTC TICCCCCOOO CAOCICCAOO GAIGIGOCCT CATIGCIGIC TOCCACICCA GAOCIGOOO ACTORCOCTO COCTOCOCAC ARCTOCTOTA CARROCAC GROTTINGTI COTOACOCAC TCAGTOCCCT CTCCAMCTAC ACCTOTGACC AAGGCTTATG TGGTCAGGAC TGAGCAGGGG TACAACITCT CACTOGIGGC CTIGOGOCIC GCATCATOGO TAATOGGAAG COCTICCAGO TOCGIGOCII CATGATITOIC ANGGGIOCO CONGCIAAN GADONCOANT CICCIGACON ACGIGNACIT CGINCICICA OTGAACTTOT ACCAAGAGOT GATGAAGCAC GCAGATCCCC GGATCCAGGG CTACCCTCTG ATOTICACTOT OGACICAGTIC TICACTICCTICC ACCICACACA CTGAAGCAGT AGCITICTIGG ACTOTOCOTO TACATTOTOT ATGAGTTCCT GATGTCGGGC 834 780 720 660 600 540 480 420 360 ĕ 180 120

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(2) INFORMATION FOR SEQ ID NO: 254:

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Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: nucleic acid (A) LENGTH: 1508 base pairs

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SEQUENCE DESCRIPTION: SEQ ID NO:

(D) TOPOLOGY: linear

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STRANDEDNESS: double

8 SS S 25 8 CCACCAACOT TOGGAGTOGA COTCATCAAT GAGOTTOTOG AGAACTTTOG CAGATOTOCO CTTCCCATGG AAGIGGICIG GICOGCAAGC CITIGICITT GICIGCCAGA CIGICATIGA GGAIGACIGC ATTICICCAT GCTCACCCIC CCAGGICAGC GAGATGGTGA AGAAGCTGCA CGCGGCAACA TIGAACITIT AAAATITIAG AICAGCAAAC ICTAAGAICC TAGAAIGGAA GCIGITCCIC GIGICTITICC TECTICCATG AGAGECGAGG TICAGIGGGC ATTEGECACG CATGIGACET AMAMICICOS AMBRIGOCAI GROCACAGOS TOOTCARCOT ACTRGRAGGO TIGRATICICO AGGGTTCCTA ACGTGCGAGT GCTGCTTGCA AAGACATTAA GACAAACTCT ACTAGAAAAA Š TOGOCTICTOC CAGCIGCCAC CAGGAGGCIG TOGAGCAGAC CATCATOGCT ACCOTOACAG COATOTCAAG TATTITIOCAA GCATCCACCC TGCCAGTACC ACCAGITIOC TOTOCATOTO ATGCCCCATO TGCTAACCTT AGCAAATGAC 254: 420 360 300 240 180 120 600 540 480 6

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	GOGATAGCTT TCGGGGGAGG AGACACCTTC CTCTCCTGCG GACTTCATTG CAGGTGCAAG	099		
	TIGCTACAC CCAATACCAG ccaatticaag aggraagaga aagtacaga aagtacaga	66		AAGG
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0	TCTTATCTTG ACTTTAAGGG GAAATAATTT CTCAGAGGAT TATAATTGTC ACCGAAGCCT	780	S	CACA
	TAMATICCTIC TOTICTICCTG ACTIGAM ACTIGAMITIG GCAGAGICALT TICCTIANGO	940		
9	AAGGAIGAG AITCCCAGAG ACCTGCATTG CITICTCCTG GITTTAITTA ACANTGACA	006	5	
2	AATGAAATTC TTACAGCCTG AAGGAGACG TGTGCCCAGA TGTGAAAGAG ACCTTCAGTA	096	2	AGCC
	TCAGCCTAA CICTICICIC CCAGGAAGA CITOCTOGGC ICTOTOGGCA GCTOTOCAGC	.1020		GACI
	CCAGCCTOT GTOTGAATCG TITGTGACOT GTOCAAATGG GAAAGGAGGG GTITTTACAT	1080	15	
	CTCCTADAGG ACCTGATGCC AACACAAGTA GGATTGACTT AAACTCTTAA GCGCAGCATA			
S	TIGCIGIACA CATTIACAGA ATGOTTGCTG AGIGICIGIG ICICAATITIT TCATGCTGGT	1200	ξ	ATA
3	CATGACCTGA AGGAAATTTA TTAGACGTAT AATGTATGTC TGGTGTTTTT AACTTGATCA	1260	3	
	TGATCAGCTC TGAGGTGCAA CTTCTTCACA TACTGTACAT ACCTGTGACC ACTCTTGGGA	1320		1
χ	GIGCIGCAGI CITINAICAI GCIGITINAA CIGINGIGGC ACAAGIICTC INGICCAAAI	1380 25	25	AICE
	AAAATTTATT AATAAGATCT ATAGAGAGAG ATATATACAC TITTGAITGT TITCTAGATG	1440		AAGAI
9	TCTACCAATA AATGCAATTT GTGACCTGTA TTAAAAAAA NTAAAAAAAC TCGAGGGGG	1500	ç	TICCI
2	CCCGGTAC	JC 1508	ર	ATGAG
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Š		35	35	ATTTA
	(2) INFORMATION FOR SEQ ID NO: 255:			ATTICO
				ATTTO
Q	(A) LEWATH: 4514 Base pairs (B) TYPE: nucleic could (C) STRANDERMECS: Amble	40	40	TAGGA
	(D) TOPOLOGY: linear			AACTC
v,	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 255:		45	TTOCT
,	GAGAGACICA CACTICTITI CCAFIAICAC TGAGGAIGTA GTGGAGATAG CAGGGGAAGA		}	ACATA
	GCACCTACCT GTGTTGGTGA GGTTTGTTGA TGAATCTCAT AACCTAAGAG AGGAATTTAT	120		AATAT
0	AGCITICCTG CCITATGAAG CCGATGCAGA AATTITTGGCT GTGAAATTTC ACACTATGAT	180 . 50	20	CCACA
	AACTGAGAAG TGGGGATTAA ATATGGAGTA TTGTGGTGGC CAGGCTTACA TTGACTCTAG	240		TACAAC
<b>v</b>	TOGATITICI TCCAAAATGA AAGTIGTTGC TICTAGACTT TTAGAGAAAT ATCCCCAAGC	300	¥	THOTE
,	TATCTACACA CTCTGCTCTT CCTGTGCCTT AAATATGTGG TTGGCAAAAT CAGTACCTGT	360	3	TTCCA
	TATGGGAGTA TCTGTTGCAT TAGGAACAAT TGAGGAAGTT TGTTCTTTTT TCCATCGATC	420		GCAGG
0	ACCACACTG CITTIAGAAC FIGACAACGF AATTICTGTF CITTITICAGA ACAGTAAAGA	480	9	TGACA

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840 1560 90 960 1020 1080 1140 1200 1260 1320 1380 1440 1500 1620 1680 1740 1800 2160 1860 1920 1980 2040 2280 2100 2220 ATTITA GIGGAACTCC IGCAAGCACT IGITITAIGI ITAGAIGGIA TAAATAGIG AITIT GAITICAITG ITACIAITGI IGITCITAAA AAIGICCIAI CITITIACAAG BOTANA GANCTIGANGS ANATICTICCE TICTICAGTOS ACAGGCAGGC ATGATGCTT NATATT AGATGGAATA ACTATATAGC TGGCCGAGCA TTTGTACTCT GCAGTGCAG KAGTA CTGCATTCAC TCAAGGAAGT GATTGGAAAA TATTGAAGTT TATCATGAA GAAAC TGTGGAAAAT ACCTAAGAGA CTTTTAAAAA TAGGCTTTCT TATATTTGAT TITGG AAAACCICC AGGGCAAAC CICIGAIGIC TICITIGGGG CCGGIAGCT JITIGA GGAAGCCACA AAITIIGGCAA CCAAACTIGA TATIICAAAIG AAACTCCCT ITICOS CAGAGOTICAC CAGGOTAACT TGGAATOTICA GOTAACOTOT GAGAGOTIAC GACAC GCTGTCAGCT GAGCTTCATT GTTGGAGAAT CAAATGGAAA CACAGGGG AATOT GTATGCATTG CTGAAGGTCC TGTGTATTCT TCCTGTCATG AAGGTTCACA AGGIC AAGTAACITG GCIFTGCITA ACATAAATIT TGATATAAAA CACGACCITC ATGGT GGACACATAT ATTAAACTCT ATACAAGTAA GTCAGAGCTT CCTACAGAT SAAGA AAAAGCCGTA AGTGTATGTA GACCACTTAA TCACTAAATA TCTTTCCCTA SCAIT TAITAICACT GIGGAICICT ACTIGITGGG IGITAIGAAT ICITICAAGA NITIT GAAGAGGIGT GGGAGGAAGG AATACAITIT AIAAAATGIT GTAGTGAAGC NITGA CCITIGACIA AIAGGAGITI TAAGIAIGIT AAAAAICIAT ACTGGACAGI BAAAT TACCGGAGAA AAGCTTGTGA GCTCACCAAA CAAGGATTTC AGTGTAGATT WANTE CANTACCICG GAGGAACACC ATGCTGACAF GTATAGAAGT GACTTACCC TCAAG CTITGAAGAC CTACCTGITC ITCCAGAAGA GAACGITGAA AGTGCCAIC ITICI TGAACITAAA GAAACAAATG ACAAAGITIG AATGGAAAAG CCTGCTGTTG ATAT GITGAACACT ICTOTITCAT GGITGAGACA GAATCAGAGG CCATGGATAC GAAAC CCTAAGTGTC CCAACAGTGG AGCACATTAT TCAGGAACTT AAAGATATA COSTA TGAAAATGGA CGAAAGCSTC TTAAAGCATA TITGAGGAAC ACTITTGACA CICCA TIGAATACAT TAGCCATIGA TAATCTACCT GITTAAATGG CCCCTGTTT ITTGC OTGATCTCTG TTGATGGCAC TCTGGAATTG TTTCAGTTAA GTCATTTTAA ICTGA TITGICIGIT TITITICICI GICITITICC AIGACICITA TATACICC ATAGA GCTTCCGTCC ACCATCTATG AAGCCCTCCA CCTGCCTGAC ATCAAGTTT CAICT CGTIGCIGIT TACATICCTT TGIGGAGCCT ACAICTICCT AAGCITTIT GAACA GCACCTCAAA GCTCTTAAAT GCTTATCTCT GGTACCCTCA GTCATGG

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1380 1320 1260 1200

1860 1800 1740 1680 1620 1560 1500 484

CCGEAGAGET CACCAGGGTA ACTIGGAATC TCAGCTAACC TCTGAGAGTT ACTATAAAGA GTACTGCATT CACTCAACGA AGTGANTGGA AAATATTGAA GTTTATCATG AATTTTOGTT ATTAGATGGA ATAACTATAT AGCTGGCCGA GCATTTGTAC TCTGCAGTGC AGTGTCAGAT AAAGAACTGA AGGAAATCTG CCATTCTCAG TGGACAGGCA GGCATGATGC TTTTGAAATT GTATCTGTTG CATTAGGAAC AATTGAGGAA GTTTGTTCTT TTTTCCATCG ATCACCACAA AAGTIGGGGAT TAAATAIGGA GTATTIGTCOT GGCCAGGCTT ACATTIGTCTC CIGCCITATG ARGCCGATGC AGAAATTITIG GCTGTGAAAT TICACACTAT GATAACTGAG CAAAACATIC TIGIGIAAAT ICICITAAAC AITIGATAAA CAGCIICACA AITC AAAAATATGG AAAATATTGC TOTTATTTTT GOTGAAGAAA ATCAATTTTG TATAGTTTAT CATCTIGATI TATAAGCAAA ACCIGGAAAA CCIACAAAAT AAGIGTIGIG GITTAICIAG ACAGCACCTC AAAGCTCTTA AATGCTTATC TCTGGTACCC TCAGTCATGG GACAACTCAA AACCCTAAGT GTCCCAACAG TOGAGCACAT TATTCAGGAA CTTAAAGATA TATTCTCAGA TGAGGAAGCC ACAAATTIGG CAACCAAACT TGATATICAA ATGAAACICC CIGGGAAATT GOGAAAAACC TCCAGOOGCA AACCTCTGAT GTCTTCTTTG CGGCCGGTAG CTTGACTGCA TITGATITCA TIGITACTAT IGITGITCTI AAAAATGICC TAICITITAC AAGAGCCITT TTAGIGGAAC ICCIGCAAGC ACTIGITITA IGITTAGAIG GIATAAAIAG IGACACAAAI TCTTCCAAAA TGAAAGTIGT TGCTTCTAGA CTTTTAGAGA AATATCCCCA AGCTATCTAC TICANICIAA ATAAAAIGIG AATTITIGITT AAAGCITAGG CACAITATIT AGAGCTICCG TCCACCAICT AIGAAGCCCT CCACCIGCCI GACAICAAGI TITITICCIAA CACGCTGTCA GCTGAGCTTC ATTGTTGGAG AATCAAATGG AAACACAGGG GGAAAGATAT INFORMATION FOR SEQ ID NO: CTTCCTGTGC CTTANATATG TGGTTGGCAA AATCAGTACC TGTTATGGGA SEQUENCE CHARACTERISTICS: AACTIGACAA CGTAATTYCT GITCTTTTTC AGAACAGTAA AGAAAGGGOT TCGGAGGAAC ACCATGCTGA CATGTATAGA AGTGACTTAC CCAATCCTGA SEQUENCE DESCRIPTION: (B) TYPE: nucleic acid (A) LENGTH: 2357 base pairs TOPOLOGY: linear STRANDEDNESS: double 256: SEQ ID NO: 256 TITGIGGGGT 2514 2460 2400 600 540 480 420 360 300 240 660 180 120 8 35 3 25 20 5 8 55 8 5 2 ARGCITTIGAA GACCIACCIG TICTICCAGA AGAGAACGIT GAAAGIOCCA TOITICCTIT AACTOTOGAA AATACCTAAG AGACTTTTAA AAATAGGCTT TCTTATATTT GATATTTOGA GTCAAGTAAC TTGGCTTTGC TTAACATAAA TTTTGATATA AAACACGACC TGGATTTAAT TAAATAAAAT GIGAATITTIG TITAAAGCIT AGGCACATTA TITTITTIGIGG GGTCAAAACA ATTINIDAGO AAAACCIGGA AAACCIACAA AATAAGIGIT GIGGITTAIC TAGAAAAATA TATOTTGAAC ACTICTOTTT CATGOTTGAG ACAGAATCAG AGGCCATOGA TACTGACAAC ANTIACCOGA GAAAAGCITG TOAGCTCACC AAACAAGGAT TTCAGTGTAG CCATTGAATA CATTAGCCAT TGATAATCTA CCTGTTTIAAA TGGCCCCTGT TTGAACTCTC OTHTCHANAR CCACCANAGC OTCTTANAGC ATATTTCAGG AACACTTTCA CAGACCAMAG TOTOTATISCA TITOCIGAAGG TCCTOTOTAT TCTTCCTOTO ATGAAGGTTG AGAATGAGCG CAGGAACCAG GATGTGGGTG CGAGGCGTGC TCCTGGCTGT TGCAGATTGC TGCACCCGGG TOGAMMATAT TOCTOTTATT TITOGTGAAG AMAATCAMIT TIGTATAGIT TATTICAMIC TGATTTGTCT GITTTTTTTC TCTGTCTTTT TCCATGACTC TTATATACTG CCTCATCTTG TCTCGTTGCT GTTTACATTC CTTTGTGGAG CCTACATCTT CCTAAGCTTT TTAGCAGGTA TGACCTTTGA CTANTAGGAG TTTTAAGTAT GTTAAAAATC TATACTGGAC AGITACAAGA TITIGAAGAGG TGTGGGAGGA AGGAATACAT TITTATAAAAT GITGTAGTGA AGCCCACAAT ATTEMPTATE ACTORIGANE TETACTION GOSTOTIANG ANTICTINA AGAMATATAT AGAAAAAGCC GTAAGTGTAT GTAGACCACT TAATCACTAA ATATCTTTGC CTATAGGACT GGTGGACACA TATATTAAAC TCTATACAAG TAAGTCAGAG CTTCCTACAG ATAATTCCGA CATTTATTGG ACTITICIEGI GCAAAAAGAT GIICAAGCCI TATTITATAC ITIGCCIGCCC CITTICICITI (2) INFORMATION FOR SEQ ID NO: 257: TICTIGIGIA AATICIC ICTIGAACIT AAAGAAACAA AIGACAAAGT TIGAAITGGAA AAGCCIGCIG TIGIITCCACA TOCOTOATCT CTOTTGATGO CACTCTGGAA TTOTTTCAGT TAAGTCATTT TAGACATAGC (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: AGTGAGCTGC AGCTCTAAGA AGACCTGTTC TTTTGAATGG AGAGTAGCAT (C) STRANDEDNESS: double (D) TOPOLOGY: linear (B) TYPE: nucleic acid (A) LENGTH: 689 base pairs

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AGCICITAGI GGALAGAGCT AGAGGATATG TGCACGTACT TCCATCTCTC TCTCTGTCTC	240	ANATOGIGG CENGENCIE CIGIOGGACA TGACCCOCAT ACTGAACATG CTCATCGIGT
CGATTTTAGC CCAGCACCAC AGGGTAGGTT CCAGTTTTTC TCTCTTTCCA TAGCTGTAAG	300	TCCBCTTCCT GCSTATCATC CCCAGCATGA AGCCGATGGC CGTGGTGGCC AGTACCGTCC
OCCUTIVIO GGANGGITC TCATTCTCCT TAATCTATTA TTGGGTCAGT TTTCCTGCAT	360	1999CCTGOT GCAAAACATG CGTGCGTTTG GCGGGATCCT GGTGGTGGTC TACTACGTAT
STCCCCAGCC TCCCATCACT GCCACCACT CCCCACAGAG ATGCCCTGCT CATCCGACTG	420	TICCAICAT ISSCANCAAC TICITIVAGAS COSTCATIOT GOCTCTICCT GGAAACAGGA
OGGETTITGAC TECCACACIO TOTIACECETE TITOTOTIGAC GECCTOCTICE CAAAACETTE	480	פככושמואפה מככשמואפים מופפפאפכרד כמאכלאפרדה פאנואמרוסים
AGCAAACAGC TITCCAAATG GAAGTIGTCA CTGTCARGGS CTTTACAATC AGCAACAGCA	540	CCAACAACTT CGATGACTTT GCGGCTGCCC TGGTCACTCT GTGGAACTTG ATGGTGGTGA
AAATCTACAT GCTGCTGAGG GTCCTGCCTC ATTAAGATGC AATAAATATG TAAGTACATA		ACAACTISSCA GGIGTITICTIS GAIGCATATC GSCGCTACTA AGGCCCGTGG TCCAAGATCT
AAAACAOCAA TAGAAGAAAC GTAATGCTTT ATTCTCAAAT ATGNATGTCT ACATAGAAAA	099	ATTITIONAL CIDENOCCIO GIOLOGICIO TCALCIGOGO CAACCIGITI CIGGCCCTGA
GCCAAAATTA TTAAGAATTAG TAAGGAATT	689	TTCTGGAGAA CTTCCTTCAC AAGTGGGACC CCCCCAGCCA CCTGCAGCCC CTTGCTGGGA
	07	CCCCAGAGGC CACCTACCAG ATCACTGTGG AGCTCCTGTT CAGGGATATT CTGGAGGAGC
(2) INFORMATION FOR SEQ ID NO: 258:		COSSOBIGEA TEACCTCACA GAGAGCTEA GOCAGCACCC GCACCTGTGG CTGTGCAGGT
(i) SEQUENCE CHARACTERISTICS:	25	GACOTECOGG TETGECATEC CAGEAGGGC GGCAGGAGAG AGAGGCTGGC ATAACACAGG
(A) LENGTH: 2377 base pairs (B) TYPE: nucleic acid		TOCCCATCAT GOLAGAGGG GCCATGCTOT GSCCAGCCAG GCAGGAAGAG ACCTTTCCTC
(C) STRANDERNESS: double (D) TOPOLCGY: linear	ç	TGACCGACCA CTAACCTGGG GACAGGAACC AAGTCCTTTG CGTGTGGCCC AACAACCATT
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 258:	On The State of th	TACAGAACAG CTGCTGGTGC TTCAGGGAGG CGCCGTGCCC TCCGCTTTCT TTTATAGCTG
TOGACCOACG COTCOSCOGA TOTGATGATT CCTOCOTATT CCAAGAACCG GCCCTATGCC	09	CTICAGIGAG AMTICCCTIG TCGACTCCAC AGGGACCTITI CAGACAAAAA TGCAAGAAGC
AICTICTICA TAGICTICAC TOTICATAGGG GACGCCCCG GCGCTGTGCT ATCCTGTGCC	120 35	AGGGCCTCC CCTGTCCCCT GCAGCTTCGG TGGTGCCTTT GCTGCCGGCA GCCCTTGGGG
GOCCACCCTT GOSTTGSTTT TGCTGCTGTA CTGGTGGCGC CCCTGACCGT GGCTGTCTCC	180	ACCACAGGCC TGACCAGGGC CTGCACAGGT TAACCGTCAG ACTTCCGGGG CATTCAGCTG
TETTGAAGGA ACCTOTITE TGATGAACET GCTGACAGCE ATCATCTACA GTCAGITCCG	240	GGAATGATAC TAATACCTCC GATTTTAGCC CAGCACCACA GGGTACGTTC CAGTTTTTAT
GOSCTACCTG ATGAMATETE TECHGACCTE GETOTTTEGG AGGOGGGTGG GAACECGGCT	300	TTCTTTCCAT AGCIGIAAGG CCCTTTCTGG GAATGGTTAT CATTCTCCTT AATCTATTAT
GCCTTTGAMG TCCTATCCTC CATGCTGGGG GAGGGAGGAG CCTTCCCTCA GGCAGTTGGG	360	TOGGTCAGTT TTCCTGCATG TCCCCAGCGT CCCATCACTG CCACCCACTC CCCACAGAGA
отваляется наласттвет вслостет смаласетсе лествелела стесслелал	420 45	TECCCTECTC ATCCERCTEG GGCTTTGACT CCCACACTGT GTACCCCTCT TGTGTGGACG
CAGGCCATGA TOGAGAAGGT GCOTTCCTAT GSCAGTOTTC TGCTCTCAGC TGAGGAGTTT	480	CCCTGCTGCC AAAACCTTCA GCAAACAGCT TTCCAAATGG AAGTTGTCAC TGTCAGGGCC
CAGAAGCTCT TCAACGAGCT TCACAGAAGT GTGGTTAAAG AGCACCCGCC GAGGCCCGAG	540	TTTACANTCA GCAACAGCAA AATCTACATG CTGCTGAGGG TCCTGCCTCA TTAAGATGCA
TACCAGTOTO GETTTOTIGGA GAGGANOCCA GETTOCITOTO GGOCAGINAC TACTITICACT	009	atabatrict archicatar aaaaaaaaa aaaaaa
ACCTIGAGGIA CETCATGGC CTIGGGIAACC TIGGTOTCCAT TTGCGTGTTC CTIGGTGCTGG	099	
ATECAGATET TOCTOCCTGC TCAGCOTGAT GACTTCATCC TGGGGGGTCT CAACTGGGTC	720 55	(2) INPORMATION FOR SEC ID NO: 259:
ITCAITIGIGT ACTACCIGIT GGAGAIGCIG GCICAAGGIC ITITIGCCCIG GGGCCIGGA	780	111 CENTERINE MIXEN MEDICATIC.
ROGIACHKIT CCTAACCCCA RCAAMGTGTT TTGAACGGGC TCCTCAMCGT TTGTCCTGGC	840 60	(1) SEQUENCE CHARACTERISTICS. (A) LENGTH: 1193 base pairs (B) TYPE: nucleic acid

2220 2280 2340 2377

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(C) STRANDEDNESS: double (D) TOPOLOGY: linear

3 6 3မွ 25 8 5 5 55 50 GAATTITIGAT CITTAGAGAC ACTAGITITIG CCCAACITAA CATITITACOT TAATTITITAC AATAAACTTT CACAGTCCAG TATCCAACAG GAACTGTGTG TGTCTTAAGA CCGAAGTTCA ACAGACCACG TGAAAGGGAA TGCTGGTCTA GCTGGCGTGG TATGTTTATA GGCGAATTTC 3 CANCITECTI GOGATOTOGO CITITIGGAA GGAAAAAAAT INCCCCAAAG GCAAATCCCA CATTAGICAT CAACATTACA IGITTICATGC TICAGATATI TIACIGCIIG IGICCIIATI GICTITITIT TAAITAATAT GIGIGCATIG TTACAANGIA TGIIGGAIGI CITTIGACCC COCCTITIGI TAAAACIGAA GAITITIGGAA AAIGGITIGIC ACIGCICTIC CAGCCTAIGA ATAGTATTIG ACACICATGC AAAATAATGT GAAAACATCT AGATTTAGTA GITTATICIG GTTTGACAAG CATTAGTGAC AAAGGCAGAA AAGATTTATC AGCCATGCTA AAAGAGTGAA ATATOCIATT TITOCIACIG TETTECTICA GEAGIGEATA TICITITIGEA AAGITEITTIG GTTGGACAGC TTTAAACAGA GTTGATGGTA CTTCAAATAT AGCTCATTGA TACTTAAGGG CCCTTTCCTT GITTIAGAIT TACTTIGCTC TICGITAATC TIATICCTGA IGATCIAGAA TAAANGCTIT TITTOTTANC AGAGATIGNG TACTATTITT ATTITTAATA AANGTANCTI titaticaaa aacaatotot teateaaagt aattoeteae attotoeagt aetatottot ATATTYTTOT GAAATOGAAC CATOGATTTA TOTCTOGATC ATCCATACAG AACCAACAAT GCCAAACTTT CCCCCAANGC TTCGAAACTT GCAAGCCGAA ACCTTGAATC GTTAAAAGTT 5 ACCAGAAGGA AGCCAAAATA GITTTTTCCT TTTGAAAAGIT TTTTAAAAAT TAITTCATGG GOGITIOCONO GGOGOCOTIOG COCGAAGAAG CGCAAITIGGC GITICCGCGAA CGITIGGCCCI INFORMATION FOR SEQ ID NO: Ξ (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 261: SEQUENCE CHARACTERISTICS: € ٤ ) TYPE: nucleic acid
) STRANDEDNESS: double
)) TOPOLOGY: linear LENGTH: 1179 base pairs 261:

1260

1200

1140 1080 1020

960 900 840 780 720 660 600 540

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GCAAAAGATT AAAGITGAAT TITACAGITA AAAAAAAAAA AAAAAAAAAA AAA ATRIBITICAT ACATGAATAT ATCCACCCAC CTAGATTITA AGCAGTAAAT AAAACATTIC

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(C) (E) ٤

N) LENGTH: 1262 base pairs

TYPE: nucleic acid

STRANDEDNESS: double

TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 260:

8

GAAAAACCCA AAGATGCAGA CAATCTCTTT GAACATGAAT TGGGGGCTCT CAATATGGCT

69

8

CAACOGCICO GCAGCCAGCC ATGICCTGCA CCCAGGACAG CGGCCCTGGG CTACAAGGAC

180 120 6 50

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 260:

4

1140 1193

1080 1020 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120 60

TOGTOGCAGE GGCTTGCTCT TGTCTTTTTC TTTTCTTTTT AACTAAGAAT GGGGCTGTTG ATOCCHACAG ACAGOCCACT CTTTGGTCAG CCTGCTGACA AATTTAAGTG CTGGTACCTG

3

TOWARDINGT CITGARCTIGA ARCICACIDO AGROCITORAS GGROCITOCCA TOTOCORTOR

ATTGAAGCAC TGCTAAACCT TCCTAGAAAC CCTTCAGTAA TAGATAAACA AGACAAGGAC

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GCGTGGCTGG TGGCTGGTGG CATAATTGGA GCCTTGCTGG

OCACTCCTOT AGGAGGCCTG TITAGGATAA ACGTAGGCCT

AAGCCATTIT GTAATTGCAG GAGCTGTCAC GGGAAGTCTT

CTGATGGCAT TTCAGAAGTA CTCTGGTGAG ACTGTTCAGG AAAGAAAACA GAAGGATCGA

AAGGCACTCC ATGAGCTAAA ACTGGAAGAG TGGAAAGGCA GACTACAAGT TACTGAGCAC

AAATTGAAAG TAGTTTACAG GAAGATGAAC CTGAGAATGA TGCTAAGAAA

20

GCTGTTTGGC AAAGACACAG TGAACACTAG TCTGAATGTA TACCGAAATA AAGATGCCTT GOSCITICCC TACGICCCAG AGCCCTATTA CCCGGAATCT GGATGGGACC GCCTCCGGA COGAGICITI GCIOCCGAAG CIGIGACIGC CGAITCOGAA GICCITGAGG AGCGICAGAA

5

CATGGAGGTG CCGCCACCGG CACCGCGGAG CTTTCTCTGT

AGAGCATTOT GCCTATTTCC

GCACOTOCOC GAGGMYTTGA AGTOCTGAGO GCTCAAGTTT GTCCGTAGTC GAGAGAAGGO AGAGGGTTAA CCTGGGTCAA ATGCACGGAT TCTCACCTCG TACAGTTACG CTCTCCCGG ATTICCAAGG ACICCAAAGC GAGGCCGGG ACIGAAGGIG IGGGIGICGA GCCCICIGGC

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S

TOTANTOGOO GIOGOCOCGO COCTIGGOOTT TIGOCOGGING GGOGGANOTI COTGIGIOTOGI

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 259:

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PCT/US98/04493

WO 98/39448

GCATTACTAC GAAAAGAAGA AAGAGCAAGT CTTCTTAGTA ATCTTGGCCC ATGTTGTAAG

ACCATAAAAC AAGCTTACAC GAGTSCTCCA ATGGTAGACA ATGAATTACT TCGATTGAGT OCCUTTOTICCT TCAGACCOGGA TTCTGCAATT CGAAAGCAGC TTGTTAAAAA TGAGAAGCOC

240 180 120

CTTCGGTTAT TTAAGCGGAA GACTACTTGC CATGCTCCAG GACATGAAAA GACTGAAGAI

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	CINGACCICA TOTTOCHECO CYCACTIONS CERTIFICADA PORTORA CONTINUADA		CCGATGGAGT CITATATCCC TGTCAAACAA CAGTGGCAAA AATGTGGAAC TGAAATTTGT
	WANTED AND THE PROPERTY OF THE		GGATTCCCTC CGGAGGCAGT TTGAATTCAG TGTAGATTCT TTTCAAATCA AATTAGACTC
<b>v</b>	COTCOTOCTG GACTICCTOT TOGACTICTT ACCCGAGGGG GTGAACAAAG AGAAGATCAC	300	ICTICHECTC TITIANGAAT GITCAGAGAA CCCAATGACT GAGACATITIC ACCCCACAAT
•	ACCACTICACG CTCAAGGAAG CTTATIGTIGCA GAAAATGGTT AAAGTIGTIGCA ATGACTICTGA	360	NATIONAL DESCRIPTION OF STATEMENT CANDESCRIPT CANDESCRIPT CTRACCAGGIN
	CCGATGGAGT CITATATCCC TOTCAACAA CAGTGGCAAA AATGTGGAAC TGAAATTTGT	420	CHARLOCKS OF STORYS ACCESSANCE ACCESSANCE ACCESSANCE ACCESSANCE
0	GGAITCCCTC CGGAGGCAGI TIGAAITCAG IGIAGAITCI TITCAAAICA AAITAGACTC	480	
	TUTTUTGOTC TITTATGART GITCAGAGAA CCCAATGACT GAGACAITTC ACCCCACAAT	540	THY CAGAGAAAC TEGAGGET TEGAGACEA CAGAGAAAC TEGAGTECTA
<u> </u>	ANTOGGGGAG ACCOTCTATG CCGATTTCCA GGAGCCTTT GATCACCTTT GTAACAAGAT	\$ 009	
3	CATTOCCACC AGGIACCCAG AGGIAATCCG AGGGGGGC CTCCTTANGT ACTGCAACCT	099	
	CTTGGTGAGG GGCTTTAGGC CCGCCTCTGA TGAAATCAAG ACCCTTCAAA GGTATATGTG	720	CCTIMICACE ANGENISCETA TOCCESCITETT ACCTICACEA AANGICATIC CTAANGICGE
20	TTCCAGGTTT TTCATCGACT TCTCAGACAT TGGAGAGCAG CAGAGAAAAC TGGAGTCCTA	780	
	ITTIGCAGAAC CACTITIGITGG GATTIGGAAGA CCGCAAGTAT GAGTATCTCA TGACCCTTCA	840	CINCATTOCA CAGGINOAGO CAGINITICAC GTOCCAGCAA CAGACCTACT CCACTTGGCT
25	TOGROTOOTA AATGAGAGCA CACTOTOCCT GATGGACAT GAAAGAAGAC AGACTTTAAA	900	
}	CCITATCACC ATGCTGGCTA TCCGGGTGTT AGCTGACCAA AATGTCATTC CTAATGTGGC	096	
	TAANGICACT TECTATACC AGCCAGCCCC CPATGIAGCA GANGCCAACT TIAGCAANTA	1020	
30	CTACATTGCA CAGGITCAC CAGTATTCAC GTGCCAGCAA CAGACCTACT CCACTTGGCT	1060 30	
	ACCCTGCAAT TAAGAATCAT TTAAAAATGT CCTGTGGGGA AGCCATTTCA GACAAGACAG	1140	(2) INFORMATION FOR SEQ ID NO: 263:
35	grgagaaaa aaaaaaaa aaaaaaaa aaaaagagc	35	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 715 base pairs (B) TYPE: nucleic acid (C) STRANEDNESS: double
6	(2) INFORMATION FOR SEQ ID NO: 262:	40	(xi) Sequen
	(1) SEQUENCE CHARACTERISTICS:	•	· DASSECTION ACCOUNTS OF CONTROL
	(B) TYPE: nucleic acid		
Ý	(C) STRANDEDNESS: double	45	AAGCAGATAT TTGAAGAGAA CAGAGAGACT CTGAAGTTCT ACCTGCGGAT CATACTGGGG
£	(D) TOPOLOGY: linear	f	GCCAATGCCA TITACTGCCT TOTGACOTTG STCTTTITT ACTCATCTGC CTCAITTTYGG
	(xi) sequence description: seq id no: 262:		GCTGGTTGG CCTTGGGCTT TAGTCTGGCA GTGTATGGGG CCAGCTACCA CTCTATGAGC
20	GGCAAACITT CCCCCAANGC TTCGAAACIT GCAAGCCGAA ACCTTGAATC GTTAAAAGTT	09	
	GGSTITGCGINC GGCGCCCTCG CCCGAAGAAG CGCAATTGGC GTTCCGCGAA CGTTGGCCCT	120	
	CAACGGCTCG GCAGCCAGCC ATGTCCTGCA CCCAGGACAG CGGCCCTGGG CTACAAGGAC	180	CAGGICTICA GOTOCTICE TICTARGIC TEGRICOTICS GOTOCTICA GOT
25	CTGGACCTCA TCTTCCTGCG CCGACCTGCG CGGGGAAGGG GAGTTTCAGA CTGTGAAGGA	240 55	
	COTOGRACIO GACTOCCTOT TOGACTICTT ACCCGAGGG GTGAACAAAG AGAAGATCAC	300	CONCONCING ACCACANTGA GAMAGGCAG COCCOSACAGG AGCGCCCCCC GATGAAGCCG
9	ACCACTCAGG CTCAAGGAAG CTTATGTGCA GAAAATGGTT AAGTGTGCA ATGACTCTGA	09	-

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 265:

8

Ξ SEQUENCE CHARACTERISTICS: 9999 () LENGTH: 1638 base pairs
3) TYPE: nucleic acid
C) STRANDEDNESS: double
D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 265:

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2 CTCCAGOGOC CAAAAGCAGT CTGAGGTATT GGGTATACTT ATACTCTATA GGGTCGTTGA

> 783 780

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GAGGAATTGT GCTTTCACCA

GAATTTCTAA GGATTTCTGG

CAACATAGCG TOCTOTGATT

CATATATTTG CAGTATGAAC TATTGCCTCT

COCACCITICT TTTGACATGG

1638 1620 1560 TACATTCAAC TCTGATCCCG

GGGCCTTAGG

1500 1440 1380 1320

TOGECETOGG TOGETETOTE AGGGTGCACA GCCCCTCATG CETOGAGCAA TGAGGGTETA CCGACAGGAG COGCOGCAGA TGAAGCGGTT ATAGCCATTG ACGATTTKGC SACINROCCAC

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TCARGCAGGA CCCTARGATG ARGCTGAGCT TTTGATGCCA GGTGCAATCT ACTGGAAATG

ATTATICAAA AAATCATOTI TATTITGAOT CCTAGGACTI AAAATTAGIC TITIGTAATA

TAGAGCTTTT TAATAGCACT AACCAATGCC TITTTAGATG TATTTTTGAT GTATATATCT

ATTOAACAAA AATOTTTCCA CTGGCTTTTG CCTGTAAGAA AAAAAATGTA CCCGAGCACA

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AGCCTAAGAA TATGATCAGG TAACTTTCGA CCGACTTTCC CCAAGAGAAA ATTCCTAGAA

TAGCACTTAC GTAAAACATT

TGTTTCCCCC ACAGTTTTAA TAAGAACAGA TCAGGAATTC

1260 1200 1140

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TARATARATT TCCCAGTTAA AGATTATIGT GACTTCACTG TATATARACA TATTTTTATA

AGACTGATTG GAATTCTTTC TOTTGAAAAG AGGGGACACC TGTACATTCT TCCATCRTCA

CACACACAAT AAAGAACCCC CTGTAAAGAC AAATAAATGA 35

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CCCCTGGTTC ACTOCAGACA GTOGCACCCC AGCACCAGAG CACAATGAGA AACGGCAGCG CTTCTGGCTC CAGGCCGGGC CCTTTACCTC CTGTGGGTGA ATGTGCTGGG 600

540

GGATGTGATC CTACTGACAG CCATCGTGCA GGTGCTCAGC TGCTTCTCTC TCTATGTCTG 480

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GCAGGGCATG GCAGAGTGAG TGTCCCCCAC CGCCAGCCCA GGCACCTTAA

OCTOGATOGO ACGAGCAGCG TICICIGAGG ATGGGGCCCT GATOGATOGT GGCATGGACC

TOGGCCTGGT TOGCCTGGGC

30

TTTAGICIGG CAGIGIAIGG GGCCAGCIAC CACICIAIGA 360 300 240

23 ANGANGCAGA TATTIGANGA GANCAGAGAG ACTCTGANGT GOGGCCAATG CCATTTACTG CCTTOTGACG TIGGICTICT TITACTCATC TOCCTCATTT TCTACCTGCG GATCATACTG

AAGTOCATOA GCTOCCGATO TOGTOCTTAG TOATTOCGGT TICCOTCGCT CICCCGTGTT TECEGOGETO GETATTIBEE TEGENECATO GEOCECANGO GENANGIGOS ENCONGRAGOS 180 120 6

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TAGTGCATCA GATGTAACAG AACAAATTAT AAAAACCATG GAACTACCCA AAGGTCTTCA CCCACAACGA ATGAATGAAC AGCCACGTCA GCTTTTCTGG GAGAAGAGGC TACAAGGACT AATTITICAAA CAACCGGTAA CCAAAGTCAC AAATCATCCT AGTAATAAAG TGAAATCAGA

AGGAGITIGGT CCAGGTAGCA ATGATGAGAC CCTTTTATCT GCTGTTGCCA GTGCTTTGCA

660

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480 420

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GCCTCAGITG GCAAGGTACC TGGGAAATAC TGTTGATCTC AGCAGTTTTG ACTTCAGAAC

AAGTGCTGGC AAGAGCGATG TCTACTACTT CAGTCCAAGT GGTAAGAAGT TCAGAAGCAA

240 180 120

GGATTIGCCCG GCCCTCCCCC CCGGATGGAA GAAGGAAGGAA GTGATCCGAA AATCTGGGCT

OTCOGOGAGO GOGOGOYCOG OGYCCAGGOG ANCCCOOGIAC ACOGIAGAGCO GGAAGAGGAT GOCIACGRAGOC GOCOGCIAGOG GTGOCOGOGGG COCOCOCOGG COGGRAGOCOT INCCCTTTTCCC

60

CARTCAAAAT AAGGGTAAAC CAGACTTGAA TACAACATTG CCAATTAGAC AAACAGCATC

TOGAMAGATG ATOCCTAGTA AATTACAGAA GAACAAACAG AGACTGCGAA ACGATCCTCT

360 300

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CATCTTOTOG CGAGCTOCTO ATACAGAAGA GATOGATATT GAAATGGACA GTGGAGATGA GAAACAGGAA GAGCGAGTAC AGCAAGTACG CAAGAAATTG GAAGAAGCAC TGATGGCAGA TIGGCITAAC ACATCICAAC COCTCIGCAA AGCTTTTATT GTCACAGATG AAGACATCAG CACAMBETET GEGECAATEA CABGGEAAGT CTEEGETGET GTGGAAAAAGA ACCETGETGT

840

780 720 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:

20

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 783 base pairs

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(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 264:

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> > S

TIGGGTATAC TIATACICTA TAGGGTCGTT GAATAAATGG CTTAGAATGT GAAAAAAAA CAGCCCCTCA TOCCTOGAGC AATGAGGGTC TAGTCCAGGG GCCAAAAGCA GTCTGAGGTA

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	493		494
	(2) INPORMATION FOR SEQ ID NO: 266:		
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1455 base bairs		(2) INPORMATION FOR SEQ ID NO: 267:
2	(B) TYPE: nucleic acid	5	
	(C) STRANDEDNESS: double (D) TOPOLOGY: linear		
10	(xi) Sequence description: Seq id no: 266;	01	(C) STRANDELNESS: double (D) TOPOLOGY: linear
•	COTOCOTACT OCCATIGNAGO TACCOSOSTICC GGIANTICOCIA GGOTICIACOC ACGOSTICOSC	09	Namods ()x)
	TCAGTTGGCA AGGTACCTGG GAAATACTGT TGATCTCAGC AGTTTTGACT TCAGAACTGG	120	The last way to the state of th
15	AAAGAIGAIG CCTAGTAAAT TACAGAAGAA CAAACAGAGA CTGCGAAACG ATCCTCTAA	180 15	
	TCAAAATAAG GGTAAACCAG ACTTGAATAC AACATTGCCA ATTAGACAAA CACCATCAAT	240	CTGCCTGTCT ACATCAGCCT GGGCTGCAGC GCGCTGCCGC CGCGGGGCCG GCAGCTGAAC
			TATGRECICT TCAGGGGGG CACCOTGTTG CATTCATCTT TGTACCCCCA GCATCTACCA
20	TTICAMALAA COMFTANCA AMBICALMAA TCATOCTIANT AMTAANINGA AMTCATANOCO	300 20	STOTTGGCAT GTAGTAGGCA CTCAAGAAAT GTGTGTTGAA TGAAGGATGC CTGTGACAAG
	ACAACGAATG AATGAACAGC CACGTCAGCT TTTCTGGGAG AAGAGGCTAC AAGACTTAG	360	CAAGOGACT TTAITYCTITIC CTGACCCTTG CTCCTATGAC ACACCTCCTC CTGACTGCCA
	TOCATCAGAT GTAACAGAAC AAATTATAAA AACCATGGAA CTACCCAAAG GTCTTCAAGG	420	CHETCACHIC TICAGAGCAG AACTOCHTA GAGAACCTGG ATGGGAAACA GCCATGGCCA
25	AGTITISCICCA GGTAGCAATG ATGAGACCCT TITIATCTISCT GTTTOCCAGTG CTTTOCACAC	480 25	
	AAGCTCTGCG CCAATCACAG GCCAAGTCTC CGCTGCTGTG GAAAGAACC CTGCTGTTTG	540	ACCESSAGE CITYTHERE ACCESSES TOTAL CITYTHEAD CITYTHANDER A
ç	GCTTAACACA TCTCAACCCC TCTGCAAAGC TTTTATTGTC ACAGATGAAG ACATCAGGAA	009	
3	ACAGGAAGAG GGAGTACAGC AAGTAGGCAA GAAATTTGGAA GAAGCACTGA TGGCAGACAT	099	
	CHITCHICA CHICANA CANADARA CANTARARA AND CANTARARA AND CANTARARA CANTARARARA CANTARARARA CANTARARARARA CANTARARARARARARARARARARARARARARARARARARA	200	AAGAAGCAGA AAATGAAAAG ATGAAGGCCA TCGGTGCTCG GAACTTGCTC AAATCTATAG
ý	CITOTCOCON OCIOCIONIA CAMMUNINI GONIATIUMA ALGONICALIO GAMILLANIO	36	CAAIGCAGAG AGAAGCTCAA CAGCAGCAAC TYCAAGCCCT AATAGCAGAA AAGAAAATGC
ç	CINAGANINI GAICHGSTAA CITICGACCG ACTIICCCCA AGAGAAATT CCINGAAAIT		ACCTAGADAG GTATCCOGTT GARTATGAGG CTTTGTGTAA AGTAGAAGCA GAACAAAATG
	gaacaaaat stitccacts scrittscct staagaaaa aatstacc gagcacatas	840	AATTIAITGA CCAAITIAIT ITICAGAAAT GAACIGAAAA ITICGCITIT ATAGTAGGAA
40	ACCITITIVA TAGCACTAAC CAATGCCTIT TTAGATGTAT TITTGATGTA TATATCTAIT	900	GECHAPACEA AAAAAGECT CTCAAAACCA AAAAAACCTC TGTAGCATTC CAGGGGCTTG
?	ATTCAAAAA TCATGTTTAT TTTGAGTCCT AGGACTTAAA ATTAGTCTTT TGTAATATCA	096	
	AGCAGGACCC TAAGATGAAG CTGAGCTTTT GATGCCAGGT GCAATCTACT GGAAATGTAG	1020	THE PROPERTY OF THE PROPERTY O
45	CACTHACGTA AAACAITHOF TICCCCCACA OTITHAATAA GAACAGAICA GGAATICTAA	1080 45	
	AIMAAITICC CAGITAAAGA TIAITGIGAC TICACIGIAT AIAAACAIAF ITITIAIACIT	1140	TOTALGITCI CACCICITAL GCITAGIIGO MACIMAGAN IITGIMMACI IICALCIII
;	TATTGAAAGG GGACACCTGT ACATTCTTCC ATCRTCACTG TAAAGACAAA TAAATGATTA	1200	
S .	TATTCACAGA CTGATTGGAA TTCTTTCTGT TGAAAAGCAC ACACAATAAA GAACCCTTG	30	TYGATG
	TTAGCCTTCC TCTGATTTAC ATTCAACTCT GATCCCGGGG CCTTAGGTTT GACATGGGAG	1320	
. 22	i GTGGGAGGAA GATAGCGCAT ATATTTGCAG TATGAACTAT TGCCTCTGGG ACGTTGTGAG	1380 55	(2) INFORMATION FOR SEQ ID NO: 268:
	GAATTGTGCT TICACCAGAA TITICIDAGGA TITICIDGCTT AAATAICACC TAGACCIGING	1440	(i) SEQUENCE CHARACTERISTICS:
			3
9	TARITITITY TECET	1455 60	(B) TYPE: nucleic acid (C) STRANDELNESS: double

WO 98/39448

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(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO:

20 2 5 8 35 30 25 S GOCACGOGAG CAGCCGOGCT GOTCCTGCTG CCAGCCGGGC GCCCGGAGTG GGGCGGCGA AAAAGCAAGG CTCCTAACTC ATTGGGAACA ACTGGACTAT GGAGTACAGT TTACATCTTC TCAGCTGTGG TTATTTAAAG CAGACTTACA TGTAAACCGG AATCCTCTCT GTATGATICEA ACTICACTITEA TECTIANACAE AGETTETETE CIGAGIGIAE TAATTEEEEAA AGCATTCCCT TCTTCAGTGT TCCTGTTGCT TGGACTTTAA CAAATATTAT ACATAATCTG CCAAACATGA ACGTTCGAGT TCCCCCACAGT GAAGTGAATC CAAATACCCG TGTCATGAAC GOGCATCACA GTGAGGGTGT AGTAGATAAA TTCAAGGAAA TAAGAGATTT GTAAGAAACT CACTOTTTCT CTGATAATOT GAAATGAGAA GTATTTACAT TOGAGGGCCA ATGGCTGGTC GODATGTACG TATTITIGCA TOCAGTGAAA GGAACACCTT TCGAAACTCC TGACCAGGGT AGCCGGGGTA TGTGGCTGAC ATATGCATTG GGAGTTGGCT TGCTTCATAT TGTCTTACTC ATTAAAGATT ATTTTTAITA CCGTAAAAAA AAAAAAAAAA AAA AGGACCAGCT TAACTTATAA TGAATGGGCA TTGTGTTAAG AAAAGAACAT TTCCAGTCAT GCTACATTTC CTTCAAGTGC actgaaaaa aattitacag ctactgaatt tcttataagg aaggagtggt tagtaaactg AATGCCACAA CTACATGGTG TICGGATCTT TGGAATTAAT AAGTATTGAA ATGTTTTGAA ACGGAAGTTT TICACAATIT CICCAATAAT ICIATATITI CIGGCAAGIT ICIATACGAA TGAAGAGGG CTTTATAAGC AGGCTGGGCA GGCCCAGCTT ATAAGTTAAA TOTTTICAAG TOCAGATTIC CATTAAATGA TOCCTCTOTT TAATACACCT ATACAAGTTI 1003 960 900 840 780 720 660 8 540 480 420 360 300 240 180 120 8

2 INFORMATION FOR SEQ ID NO: 269:

E SEQUENCE CHARACTERISTICS: LENGTH: 1234 base pairs

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€ G TYPE: nucleic acid
STRANDEDNESS: double TOPOLOGY: linear

55 50 8 ACATATOCAT TOSSAGTIOS CTTOCTTCAT ATTOTCTTAC TCASCATTCC CTTCTTCAST CCAGTTOTAT CAGTGTTGAT TCATTTCATT ACTTCCTACA GAGCAAACAT GAACGTTGGA GAAAAAATAA TCCTCGCAAC ACAGGTACCT TGTCATOTCA GAATTOGGGG TGTTAGGTTG ATCAGCATCT ACAAGTAGCA TATTITIGGAT GGTGTTTIGTG TGCTACTICA AAGTAACTAG GTTGCCCACA GTGAAGTGAA TCCAAATACC CGTGTCATGA ACAGCCGGGG TATGTGGCTG ž SEQUENCE DESCRIPTION: SEQ ID NO: 269: 300 240 180 120 8

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35 9 INFORMATION FOR SEQ ID NO: 270:

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CASTICCCAC TRANSAGGER ACTITITITIGG TITTICCTICGG CITRATIATIG TGTATIOGIC

AATGAGGCCA TITTITACANT TATTAACGIT ACAG

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GCAGACTTAC

AUGUADACCO GARICCICIC TATACAAGIT TATTAAAGAT TATTITTATI ATTOTOTTAA GAAAAGAACA TITICCAGTCA TICAGCTOTG GITATITIAAA

TTTCKCTTGT

TITATGTAAG YOGATGTATA TCCTCTTGTT

TTATACAAGC

1200 1140 1080 1020

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GCTTTATAAG CARGCTGGGC

ATTICAAGGAA ATAAGAGATT TOTAAGAAAC TAGGACCAGC TTAACTTATA

960 900

AGGCCCAGCT TATAAGTTAA AGGGCATCAC

ACTGAGGGTG CTGAAGAGGG

GIGCAGATIT CCATTAAAIG AIGCCICTGT TIAATACACC

TGGTACATTT

840 780

TOAAATGAGA AGTATITACA TIGGAGGGCC AATGGCIGGT CCITCAAGIG CIGITITGAA

GCTACTGAAT TTCTTATAAG GAAGGAGTGG TTAGTAAACT GCACTGTTTC

TSTGATAATO

720 660 8

GTICGGATCT TIGGAATTAA TAAGTATIGA AATGTTTIGA AACIGAAAAA AAATTITACA ATCCTAAACA CAGCTTCTCT CCTGAGTGTA CTAATTCCCA AAATGCCACA ACTACATGGT TCTCCAATAA TICTATATIT TCTGGCAAGT TTCTATACGA AGTATGATCC

AACTCACTTC

540

480 420 360

CATTOGGAAC AACTGGACTA TOGAGTACAG TITTACATCTT CACGGAAGTT TITTCACAATT CATOCAGTGA AAGGAACACC TTTCGAAAACT CCTGACCAGG GTAAAGCAAG GCTCCTAACT GTICCIGTIG CTIGGACTIT AACAAATATI ATACATAATC TGGGGATGTA CGTATITITTO

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8 Ξ SEQUENCE CHARACTERISTICS: <u>0</u> (B) TYPE: nucleic acid (A) LENGTH: 574 base pairs STRANDEDNESS: double

(D) TOPOLOGY: linear

CACTOCTACC TOGTACTOCT TTCAGTOTOT TCCCCCTCAG CCCTCCGGCG TGTCAGGCAT AAAACGTCAG TCTGCCTGTA AATTTCAGCA AGCCGTGTTA GATGGGGAGC GTGGAACGTC TATOGRICCC CATGRAGCCC TACTACACCA ARGITTACCA GGAGATITOG ATAGGAATOG ACTOTACACT TOTATAAGTA CCGTTTACTT CATGGCATGA ATAAATGGAT CTGTGAGATG AAGCTICAGC OCCIGCICCT GGICATCACT AACCAGATIT ACTICGAGIA CATGIGAAAG GGCTGATGGG CTTCATCGTT TATAAAATCC GGGCTGCTGA TAAAAGAAGT AAGGCTTTGA NGAGGTOCGT TCTGAGCCGT CTGTCCTGCG CCAAGATGCT TCAAAGTATT ATTAAAAACA ž SEQUENCE DESCRIPTION: SEQ ID NO: 270: 420 360 300 240 180 120 60

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ACTOTGAGTA GATAATTIGT CATGCAGGGC ATGCAATCAG AATCTCACTG AGCCACCAT CATTOTGAAA TAATTACCTC AGTTGTACAG GACTTGGTGA TCAGGATCCA GGCACTCACT

TOTATTICTAC TOCTCAATAA ACCITITATTA AACT

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1380 1320

1440

1500 1560 1620

1680 1731

480		CTTTTAACTS AAAAGGANTG GANTAGAAGG GTTTGCAATG CCATATTATT. GGTGGAGGGC
540	v	TOTITITAACA TCTTTGAAGT ATGGCTTGCT GAATATCTTT ACCAACATCT TGAATATATA
574	3	TICTAGISTIC CACAAGAITT AGCAAAAGA TAAAGCTIGG GIGGAATATC AITITAAAAJI
		GITCATOTIC TOTTCTATAT TITCTTCACC TACTCTCCAA ATATTGTAAT GCAAAAAGTC
	9	TCAGRANGA TITIGGRAGIA TIPARITITIGT GGICATIGIT TCTCTTCCAT AAATITIATIT
		TCATTAAATA CITRITAGAG GGITTIGAAA TGITTITCAA ATAIGTGAAA TGIGAAACTĠ
	Ā	CIGICITITIA TATTAAAGTA ATTAAAGAAA AIGTATTGIG ATTGAAATTA TITTGACCTC
	2	CACAAGATGG CTCTATGAGT ATTCTTCCAG GGATTCTAAT ATTTATTAA G
09	20	(2) INPORMATION FOR SEC ID NO: 272:
120		(i) CENTIDATE CHARACTERISTICS:
180	25	(B) TYPE: nucleic acid (C) STRANDEUNESS: double
240		-
300	ç	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 272:
360	ર્ક	CTOCTTAGGA AGAGAAGGTC AGAGTTCGCG GGGGCAGAGG CATTCTTGCC GCTGGCCCAG
420		TCACTATGTA GTGGAGGGGC AGACACCCTC CCGCAARTTC TGGAAGGTTC TTAGTCTCGA
	35	CTAGGGCHOT AGCCCAGGAC TOCTAGTOGC CGGCTTCAGG TCACTGCCGG CTGAACGGAG
540		CTGCCGTCGC CATGTTTGGC TGCTTGGTGG CGGGGAGGCT GGTGCAAACA GCTGCACAGG
009	Ş	AAGTOGCAAA GEATAAATTT GTTTTTGACT TACCTGATTA TGAAAGTATC AACCATGTTG
099	€	TOSTITITAT GCTGGGAACA ATCCCATITC CTGAGGGAAT GGGAGGATCT GTCTACTITT
720		CTTATCCTGA TTCAAATGGA ATGCCAGTAT GGAAACTCCT AGGATTTGTC ACGAATGGGA
780	45.	AGCCAAGIGC CAICTICAAA AITICAGGIC TIAAAICIGG AGAAGGAAGC CAACAICCTI
840		THOGAGCCAT GAATATIGTC CGAACTCCAT CTGTTGCTCA GATTGGAATT TCAGTGGAAT
006.	Ş	TATTAGACAG TATGGCTCAG CAGACTCCTG TAGGTAATGC TGCTGTATCC TCAGTTGACT
096	3	CATTCACTCA GITCACACAA AAGAIGTIGG ACAAITTICIA CAAITTTGCT TCATCAITTIG
1020		CTOTCTCTCA GOCCCAGATG ACACCAAGCC CATCTGAAAT GTTCATTCCG GCAAATGTGG
1080	55	TICTOCABAT GGTAIGAGGC AINTICTGTC TCCAATATTA AGGCTTTTTA TAACTGAATA
1140		TCTATTITIGT CTATGAATAT ATTCCTTTTT TGACATTTAA ACATATTCTT TTATTGTGAA
1200	9	CATCAGCACT GCATGCCATT AAAGTATGTA CTATAGAGAT CTGATGAGAA ACAGTTCTTA
1260	3	

ATTTAACAGG TGTAAAACTG GTACTTCCAA AGACCAAGTT TTCAATGGTA TTACCAGAAG

TAGAAGCGC ATTAGCAGAG ATTCCCGGAG TCAGGAGTGT TGTATTATTT GGAGTAGAAA CTCATGTOTG CATCCAACAA ACTGCCTGG AGCTAGTTGG CCGAGGAGTC GAGGTTCACA

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THOTHGOTGA TOCCACCTCA TCAAGAAGCA TGATGGACAG GATGTTTGCC CTCGAGGGTC TOSCITCRARC CHOGGATCAT AGTGACCACG AGTGAAGGCT GITCTGCTTC AGCTGGTAGC TGATAAGGAC CATCCAAAAT TCAAGGAAAT TCAGAATCTA ATTAAGGCGA GTGCTCCAGA GTCGGGTCTG CTTTCCAAAG TATAGGACAT TTGAAGAACT GGTATGCTAC TCACTGGTGA

35

AGGACAGTCA GGTGAAGGAC TGTAAGCCCA CACAAGCTCT TCTTATCTCT ACTAGAATTA AAATGTTAAG TCAAAAACGG CTCCTTTTTT GCGCCTCCTA GTGAACTTAA CCAGCTAGAC

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CTACTGTGTA CTCTGATCGT ATCTTTCCAA AGTGCAGACT CTTGTGAAGT TTTCTTAAAT TOTICACTIT AAAGAAATG ACGIACCAAC AATGATITIGG CITITIAITATI ACTGIAAGAT GITATAAIGI TAAIGIGGAT GIAGIGCIITI TACITITACAG AITGAITGGA AIAAGAITAI TOCATATGAA TITTACCCACA GGACTICTGAA TCATGITTACC CACTCCCCTC ACAATGITGT CCACTTAGTG AGTTGCATTG ATCTATCCGT ACCAAATGAT GTTGAATAAT TACATAICTT

S

CATTIGAGIA CCAGCAITTA GITACAAACG ICAAAGGCIT CCGGIGCIGC TIACCITICCT TITITIGITIAA TGIGCITITIA ITITAITIAAAA AAAAITIACAA TGAAGATGCC TGITITIGICI

45

GCTGCAAGGT GCGCTCGTG CCGCTGCAGA TCCAGCTCAC TACCCTGGGA AATCTTACAC CITICAAGCAC TOTOTITITIC TOCTOTOATA TOCAGGAAAG GITCAGACCA GCCATCAAGI ATTITIGOGGA TATTATTAGC GIGGGACAGA GATTOTTGCA AGGGGCCCGG ATTITIAGGAA ITCCTGTTAT TOTAACAGAA CAATACCCTA AAGGTCTTGG GAGCACGGTT CAAGAAATTG

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 271:

(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1731 base pairs
(B) TYPE: nucleic acid
(C) STRANDEINESS: double
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 271:

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8 120 180

8 360 420 480 540 909 999 720 780 840 900

TCTKGACTAT ACTGATTTCT TATTTTGGTC ACTATTACTA AATCTCTGTT AATAITCTCT

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CCCTATGTTT TAGGGACCAC TATTAAAGCT TATAAATATT TGTGTATTTT CATTTAGAAG CCCTAARTAT TITGITATAT TOTCGCCATT ATGAATITAT AAAGACAGGA AAATATAGTT

1020

960

PCT/US98/04493

30 25 5 45 8 35 S 8 TACCATCTAT GAGAGTAGTT TATACTGCAC TGTGTACATG AATGGCTAAT GAATCTATTT GCCCTGATGT GGAGTACCTT GCCAGCATCT GCTGGGGTGA ACTITATTIT AGCCCTTCCC GATIOCCTAGG AAGAGTITIOT TGAGAAGIOG TACCATIGGIG TAGCATIGGGA GAGCATIGGG GGWTATATGG CCTTGGCGAG TCAGCCAGAA GGCATTCATT AAGTAAGTCC TGACTTTGTG CCCAGCTCTG TGTTATAGGC TTCCCAGCAC AGCGCTCTG GAAGAGGCAT GAGGCATTTC ITTCAGGAAA TGRTCAITAT CCCTOGAGAG GGGCTGCTGT GCCAGCTTOG GGAGGGTCTG GGATGGGGCT GCCCCTGATG 5 ATCTGCTCAT CTGANGAATG AGGAAATAGG AGTGAATTTG ATNITTCCTA GGTCCNTCTA AATGCACTAG GTTTGAAITT GGCATAATGG TAGCTATGTG ACCCTGAGCA AATTTCTCTC TOCAACTITO COGTOTITTA TAGATATTIC TITTCACTIT GAGTATOCTA GAGATGGGAG GCGTAATTAT TACAGAGGCA GTCCATGTGC ATTGT TTGTTOYTCT TATGAAGAAC AGAGGAGGG TGGGCAGGTC AGTGATGTCA GCAGTGAGTA TGACACCCAT AAGGAATTCA TGAAGAAAGT AGAAGAAAAG CGAGTGGACG TTAACTCAGC 2 INFORMATION FOR SEQ ID NO: 273: INFORMATION FOR SEQ ID NO: 274: E E E (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 274: CACACTITGA CITTGCTACC ATGGGCTGTG TCTANGNACG TATATATGCT TCCAAGAAGA CCCGGGGGCG GTGAAAACCC ACTCAGGAGG GGCARAGGAC GCTAGKTTKT AGWTAACACG GAACCTCARA SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 273: (A) LENGTH: 515 base pairs (B) TYPE: nucleic acid (A) LENGTH: 2995 base pairs TOPOLOGY: linear STRANDEDNESS: double TYPE: nucleic acid STRANDEDNESS: double TOPOLOGY: linear TOTOGACTAA TOCTCACOGO 1320 1260 1200 1080 240 515 480 420 360 ĕ 180 120 6

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ઝ 30 25 20 S S ટ 6 2 9 8 S CATTITIAGCA TOCAAGITICO COACCACOAA GITAGAGATG ACTIGOTOGIG CTGACATTITI CCAGCAGCGT CTTGAAACGG CCTTGTCAGA ACTGGTGGCT AATGCTGAGC TCCTGGAAGA ACTGAATGAT GCCTTGGATC GGCTGGAGGA GTTGAAAGAA TTTGCCAACT TTGACTTTGA CAACCAGCIT ICTOOCCOCT GGCAGCAGGT GTGGCTGTTA GCACTGGAGC GGCAAAGGAA TCAGCCAACC CCTCCTCCCA TGCCAATCCT TTCACAGTCT GAAGCAAAAA ACCCACGGAT GATGACTCGC AAACAGCCTG ACGTGGACCG GGTCACCAAG ACATACAAAA GGAAAAAACAT CCCGCAGAAC ATTGACCGAG TTAAAGCCCT TATCGCTGAG CATCAGACAT TTATGGAGGA AGTAGCCATG GGAGAAGTCA TCCTGGCTGT CTGCCACCCC GATTGCATCA CAACCATCAA CATCAGGTAG CAAGTTGAAA CGACCAACAC CAACTTTTTCA TICTAGTCGG ACATCCCTTG GOTGTACATO CATGOCATOT TOTOCAGOCA COCCAGOCAG TOGAACOAAG GITATOCCAT CATCHTCCCG GGCAGCTTCC CCTACTCGTT CCAGCTCCAG TGCTAGTCAG AGTAACCACA TACCAGAGGG ACCATCCCAG GGAATGACCC CCTTCCGCTC ACGGGGTCGA AGGTCCAAAC GCGCAACCUT GATOOTTCGC GTTGGTGGAG GATGGATGGC CTTGGATGAA TTTTTAGTGA CCGGTTCTTC CTCGGCAATC AGTTTGGGGA TTCTCAGCAG TTGCGGCTGG TCCGTATTCT CAAGGATGCG TATCGACCAA CAACCGATGC AGATAAAATC GAAGATGAGG TTACAAGACA CGACCGAGAT GGGGATGGTT ACATTGATTA TTATGAATTT GTGGCTOCTC TTCATCCCAA CTTCCGGCGC ATTGATAAGG ACCAGGATGG GAAGATAACA CGTCAGGAGT TTATCGATGG TOTOTOGROG AAAAAOTATA TOOGTTOGAT GAATCACAAA AAGTOTOGAG TGATOGATTT AGAGCCTACT CACOOGCCTT TCATAGAGAA ATCCCGCAGC GGAGGCAGGA AATCCCTAAG ACTICIOSCA TOGRISCAGI GOSCIGAGAC CACCCICATI CAGCOGGAIC AGGAGCCAAI ACACTIGGATO ACCATOATOO GAGOTOGOTT CGAGGAGGTO CTGACATIGGG CTAAGCAGCA CANACCTTCC AMMICCCAN CUNTOTCTAN GARGACCACC ACTOCCTCCC CCAGGACTCC OCCAGCAGCC GOCGAGGAAG TGACGCTTCT GACTTTGACC TCTTAGAGAC GCATTGCTTG GACCCTAAAA AGTCTGCCAG TCGCCCTGGG AGTCGGGCTG GGAGTCGAGC CGGGAGTCGA CTOGTOATAC CAGCAATTAG TICTICCCCG GCCTCCACAG GTGCCAAAAC TAATCGGGCA AAAATGATCC CTGCCGAGCA CGAGGTAGAA CTAACATTGA ACTTAGAGAG AAATTCATCC AGTOGOTICAG TOCAAATOTO CAAAAAGGTT TICAGGTOGAG CAGATIOGAG AGAATAAATA AGGICCCAAG CGATAACACT GTCTAAGCAC CCCCAAGCCA CTATCCACTT TICCGACACI TCAGAAAGCA GOGCIGCAGG GGGCCAAGGC AACICCAGGA GAGGGCIAAA TCCATACATT GOGTGTATAT TTATTCTGAA COOGAGAAGT TATATTGTTA AAAGTGTAAA TGAATCCTGC 1140 1080 1500 1260 1200 1020 1620 1560 1440 1380 1320 1860 1800 1740 1680 960 8 780 660 600 540 480 420 240 720 360 300 180 120

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AGAATAAITG TGFTARAAG CIGCCITAIT ITTITICITT ITGTAAGITA CIAITTICAI	1920	CACCACCAC ACTIVITIES CONTINUED TRADITIONED BACTICATE GRACGSCOOL
STGANTATT ATGTAGATAA AATTTGCCTC CTGGTAACCC TGTAATGGAT GGGGCCCAGA	1980	TATALOG CONTROL OF THE PROPERTY OF THE PROPERT
aatgaaatat ttgagaaaa caagtgaaaa ggtcaagata caaatgtgta ttaaaaaaa	2040	
AAAAGCCTAT TAATAGGGTT TCTGCGGGGT GCAGGGTTGT AAACCTGCTT TATCTTTTAG	2100	GACCAGATAT ACCATCTICG ATCTCCAGC TCCCCTCCAA ACTACATGTA TAATCCTATC
GATTATTCCT AAATGCATCT TCTTTATAAA CTTGACTTGC TATCTCAGCA AGATAAATTA	2160	
TATTAAAAA ATAAGAATCC TGCAGTGTTT AAGGAACTCT TTTTTTGTAA ATCAGGACA	2220	CAMPACHICAL CICCACANG GAGOTGTATG GAGANCCTGA
CCICAAITAG CAAGAACIGA GGGAAGGGCT TITICCAITG TITIAATGITI TGIGAITITIT	2280	CANAGEMENTS ATTACTIONS CONCOTTANT CONTINUES CTOMOSOCOTO CTNOGATIGNA
ACCTAAAGAG AGGGAACCTC ATCTAAGTAA CATTTGCACA TGGATACAGC AAAAGGAGTT	2340 15	
CATTGCAATA CTGTCTTTGG ATATTGTTTC AGTACTGGGT GTTTAAAGGA CAAATAGCTG	2400	CERCECATEC ADMINISTRA CANTITUESS CERCECATEC ACATGRAPIGA TOSOCCAGTA
CTAGAATTCA GOGGTAAATG TAAGTGTTCA GAAAGGTCA GAACATTTGG GGTTTTAAAC	2460	
TGATTTOTTS CTCCCTATCC AGCTAGACA CCAGTAACTC TTGTGTTCAC CAGGACCCAG	2520	
ACCCITGGCA AGGCATAGGC TCGITGGTGA CALTGTGAAT TICAGATTIG TITTATICCAC	2580	
TITITITICCI ATTITITIAA ATGGTCGATC AACTTCCCAC AAACTGAGGA ATGAAFTCCA	2640 25	
CEAGCCTOTT CTGAAAATGT GGACGTAAGA CAAACACGTG CTCGTCCTTT AATGGAGTTC	2700	CONTRACTOR BACTABAR ARCHICA CACATCAAAA AACCAAACTI CATCCTTCCC
ACCAGCACAC TIGITAACCA GICCIGITIG CITICGICIT ITITIOIGGG TAARAAGIC	2760	
AACTGACCAA GTGACCATGA AAAGGGGCTG TCTGGGGCTC CTGTTTTTTA GCTGCTGTTC	2820	
TTCACCTCCG ACCATGITGC 191916AITA TCTCAATTGG TITITAATTGA GGCAGAAACT		מואריו איר איר איר איר איר איר איר איר איר איר
GAAGCICTAC CAATGAACTG TTTAGAAACA AGACACACTT TTGTATTAAA ATTGCTTGCA	2940 35	
PHILIPPINE MAINTAINS AMARIANA ARABANTAN ACCORDIONAL		ACTICATICG ATGIATITIT GCCTTTITIT TGTTGTCGTT TAAAGAAAGA CTTTAACAGG
CINCLUMEN ACCORDING ANGERON ACCOUNTING MOSSOCIAL GOLD.	6000	TETCATERAG AACAAACTOG AATTTCATTC TGAAGCTTGC TTTAATGAAA TGGATGTGC
	40	TAAAAGCTCC CCTCAAAAAA CTGCAGATTT TGCCTTGCAC TTTTTGAATC TCTCTTTTÄ
(2) INPORMATION FOR SEQ ID NO: 275:		TGTAAAATAG CGTAGATGCA TCTCTGCGTA TTTTCAAGTT TTTTTATCTT GCTGTGAGAG
(i) SEQUENCE CHARACTERISTICS:	, z	CATATOTTOT GACTOTOTT GACACITITA ITTACIGOTT ICITIGIGAA GCIGAAAAGG
(A) LENGTH: 1990 Dage pairs (B) TYPE: nucleic acid	7	AACATTAAGC GGGACAAAAA ATGCCGATTT TATTTATAAA AGTGGGTACT TAATAAATGA
(C) STRANDENESS; double (D) TOPOLOGY: linear		GTOSTTATAC TATOCATAAA GAAAAAYCCT AGCAGTATTG TCAGGTGGTG GTGCGCCGGC
(xi) SEQUENCE DESCRIPTION: SEQ 1D NO: 275:	50	ATTGATTTTA GGGCAGATAA AAGAATTCTG TGTCAGAGCT TTATGTTTCT CTTTTAATTC
GGGACCCGCG CGSCTCCCGG GGATGGTGAG CAAGGCGCTG CTGCAMOGTG TCTGCCGTCA	09	AGAGTTTTTC CAAGGTCTAC TTTTGAGTTG CAAACTTGAC TTTGAAATAT TCCTGTTGGT
ACCOCAGAGG ATGAAGCTGC TGCTGGGCAT CGCCTTGCTG GCCTACGTCG CCTCTGTTTG	120 55	CATGATCAAG GATATTTGAA ATCACTACTG TGTTTTTGCTG CGTATCTGGG GCGGGGCAG
GOCCAACTIC GITAATATGA GOTCTATCCA GGAAAATGGT GAACTAAAAA ITGAAAACGAA	180	GTTGGGGGC ACAAAGTTAA CAFATTCTTG GTTAACCATG GTTAAATATG CTATTTTAAF
GATTCAAGAG ATGGTTGAAC CACTAAGAGA GAAAATCAGA GATTTAGAAA AAACCTTTAC	240	AAAATTTGA
CCAGAAATAC CCACCAGTAA AGTITTTATC AGAAAAGGAT CGGAAAAGAA TTTTGAWTAA	300 60	

.1980

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E SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 276:

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double (A) LENGTH: 2436 base pairs

TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO:

20 7 30 25 TITHAAGICC GITGCATIGA AAATAACAAA CAATATCAAT GITITAATCA AGGATCICTI CCCHOCTTGG GGCACACGAG GAAATCGTAG TCGGGGAAGA CTCTACTGAA TAAGACATCA CCACATTCCT CCTTCTTATA AGAGCACAGT AACACTATCC TGGAAACCTG TACAAAAGGT AACTTCGCTT AGCTCTCCAG GGTNAAACGG GTGAGNCCTT AAAAACAGAA GAGAACAAGA TIGIATOTAT GACCIACTIT IGIAACAGAC CAIGGIIGIG ICCAAGGIAA AACCACAGIG GOGGATOICC CTTEAAACAG ACTOCTOCCT TCAGCTAAAA ACTTAATGTT CTTEATACCT GCATTOTICA GCATIGICAT GAGCITAATA TACITAAATT CTACTACTCA TIGGATIGCO CAGCAATTTG GGCAACTTTA ATTATGAGCA GAGAGGAGCC TTCAGGGGAA GTAGAGGTGG TTCAGCAGGA CCAAAAAGAG ATGCCAGGCA GATTTATAAC CCTCCCAGTG GGAAATATAG TGAGATTGGG CAAAAGAGAG CCAGTGAAGA TACAACTICA GGTTCACCAC CCAAGAAATC 180 120 660 600 540 480 420 360 300 240 8

ઝ TCAAATTGTG AATCTTTTAA ACATCTTGAT AATTTGTTGT TGAGAGCTGT TCATTCTAAA ATATITITIGG AUGCITITGIC IGCAAUCTIG ACTIGITITIT GCAGTAUCAT TATICAGACT

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6 TITICCICIT CCCICITAGI TITITACCCA ATATATIGAG AAGAGTAATG GICAATCITA ATGTAATGAA ATTCAGTCTA GTTCTGCTGA TAAAGATCAT CAGTTTTGAA AGGTTACTGA 780 900 840

45 AGGAGAAAGT AAGTIGCITT GCACCGCCIA CITAATICIT TICCATATAT IGIGATACAA 1080 1020

ACTITIGAAT ATGGAATCIT ACTATITIGAA TAGAAATGIG TATGTATAAT ATACATACAT ACATAAGCAT ATATGTGTGT GIGIGIGIGT ATATATATA ATATGCATGC TGTGAAACTT

GACTACACAA CATAAATCAC TITITAAATT CCAGGAACGG GTAGTCTGAC ACGGTGATTA 1140

50

TCCTTTTGAG GCTGAATCCG TTATTAACTT GTTATTTAGG TITITIACICC CAGTAGCAAG 1320 1260 1200

ATGTACATAC AGGTTTCTAC AGGAAGAGAT GGTATAATTT ACAATTTGGA GATTTAATAA

1440

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GTCATAAAAG CAAAATACTT ACATAGCTTT CTTAAAATAT AGGAATGACA TTACATTTTT ACATITIGIT TIAAITOITI AATAAAGCIG CIGGGCAGIG GIGCAGCAIT CCIACCIAGI GGATTCTAAG TTAGITGCAC TTACATGATT ATTGTTATIT AAAACTAAGA ATAAAGGCTG CATTITICADA GATADATIGG DATIGCIGIT GOTGADATAD CANCCADART ACTGADICIG 1380

960

30 23 20 5 5 S GAACCTAAAC TCCCTTGGAA TCTGAACAAA GGAATATAAA ATTGCCATTT GAAAACTGAC GCACATGTGT ATGGGGAAAA TAGTTCTGAA ATTITIOTITI CACAGITAAT CTICCCICCC CAAGITITOCT ATTCAAATCA ACTOCCIGAA TAATGICATG GGCATTITTA GTAGCATAGA TATAAGTIAG CCAATAGAAT TITTAGGTTA AAACAACAGA TGGGGGGTTT GTGGAGTGTT CCAGGGCTAC CCAGAAAAAG TGACTTGATA ACATGGTACC AATAAGTAAG GGATGCTCTC CAGCTRATICT GGACCTCAGA GATAGATICAG CCAGTGGCCC AAAGCCATTT CAAGTACAGA GGTCAGCTTG GCTATGGAGT GGTGGCAATA GICTICATTA CCATTACCIC TACACIGCAG TGACATTTCT TOGGTTTGCT TITGCCACTT TCAAGATTIT AACTICICAG GITATTAATC AAAATTATIG GATACIGAGI IGACIGITEC CITATECETE ACCETTECCE TICCETITEC TAAGGEAATA CTITATAAGC ATATTIGIAA ACICAGAACT GAGCAGAAGI GACTITACIT TCICAAGITT TITGAAATAA ATTICCTTIT GTAATTITAA AAAAAA TITITCITIT GCAAGACACC IGITTAICAT CIIGITTAAA IGIAAAIGIC CCCIIAIGCI GTGCACAACT TAGGTTATTT TTGCTTCCGA ATTTGAATGA AAAACTTAAT GCCATGGATT AATTATAGAG ACTACAGCTA AATAAATTTG AACATTAAAT ATAATTITAC CACTTITITOT AGTAGICIGA TGTATTTTTC TGAGGAATAG TTTGTGATTC CAATGCAGGT AAGAAGCAAA ACTCCTTTAT TAGAATTACT ATCTCTAAAC ATTCCAAAAG ACCATGAGCT AGGCTAGAAT GATACAAGTG AGCAAAAGTT CCCTTTGTTC TGCATTTGAA TGTTTCGTAT . 1680 2400 1860 1800 1740 2436 2340 2280 2220 2160 2100 2040 1980 1920 1620 1560 1500

(2) INFORMATION FOR SEQ ID NO: 277:

£ SEQUENCE CHARACTERISTICS: (A) LENGTH: 782 base pairs

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(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

Ě SEQUENCE DESCRIPTION: SEQ ID NO: 277

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8 SCAGGGCARA GOTGGGAGTC AAACCCGGGT GACAGGTGGG TGGAGAGCCCC TGTTTGAGGT TCACTIONG CONTRAINA AANTCONAAG GGCCTGAAG AAGACATITC TACTGCAGAG TGACAAGTTT AGTAGTCCCA AAATOGGTTA TATCCCTTCC CCCTTTACAT CAGAATCTTG GOTTAGAGGC ACTTGAGCAA GGCCCCCACA TCCCAACTCT GGGAGTTOTG GTGGGAGGAG GCCACIGACT TCTCCCACCC TTCTGTCTCC CCCATAATAG TTTATTTGGT TGGTCTGGAC GCACTICIGG GGGATAGGAC CAGACAAGAT AACAGGAGCT CACATGGNAA GCAGAAGCTG RUAAATOGGA AAACAACAGA AGGAGGGGAT CAAAGATAGC TGATCTCACA TGCTTCCCAG 240 120 420 360 300 180 8

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<b>W</b>	WO 98/39448	771598/04493	WO 98/39448	PCT/US98/04493
	205		908	<del>-</del>
	TOTOCCIGAT CCCICTCTGG TATTACTITT TCCCCTGGGA GCAGGAAGCC CTAGGAAGAG	087	U	961
u	GRANCTICANG GOTOCCCRGG GRATCTITICC TOCCTCCCCT GCATGAGGCA GAGGCAAGCT	540		
n	OCCITOCCAAC COCCITCCCTC AAGGAATGGC CTTGCCCAGG AATGCCCAGC ACACATACCC	009	(2) INFORMATION FOR SEQ ID NO: 279: .	
	TOTTOTITIT TICINGTORA ACTOTIGIT ATTOCITISSC TISCOTOCOT COTTOCIOCO	. 099	DE CHARACTERISTICS	
10	CICTCAACCT TTACTICTGA TITICTATITIC ATGGAATTIG GGATTGAAGT TAAACTAGAA	720		
	CAGTOCCOCC AACACCAAGT CTTGCAGGAA AAAAATACAA AGAAATTTAA CAAAAAAAA	780	(D) TOPOLOGY: Linear	,
7	V	782 .	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 279:	
3			COCOCTITICS AGITCOGICT CCTGOTOTAC GOCCAACGCC AAGIAGGGGA TIGCGITCCC	09
			TCCAGTGGCA GCCCTATCAG ATTTGGATAT GTCCTTCATA TTTGATTGGA TTTACAGTGG	120
20	(2) INFORMATION FOR SEQ 1D NO: 278:	2	20 ттгоасаот стостасаот ттттаосатт ататаасаа астоставае тостаттте	180
			TGGNTTGGAT ANTGCAGGAA AAACAACATT GCTACACATG CTAAAAGATG ACAGACTTGG	240
č	(B) TYPE: nucleic acid (C) STRANDEINESS: double		ACANCATOTC CCAACATTAC ATCCCACTTC CGAAGAACTG ACCATTGCTG GCATGACGTT	300
3	(D) TOPOLOGY: Linear		J TACAACTITIT GATCIGGGIG GACAIGITCA AGCICGAAGA GIGIGGAAAA ACIACCTICC	360
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 278:		TOCTATCAAT GOCATTGTAT TICTGGTGA TIGTGCAGAC CACGAAAGGC TGTTAGAGTC	420
30	GAGITICOGGC TOGRAACCOG TOCTCTGGGC CGGCGCTTC ACCATGGCCT CGGCAGAGCT	30	0 лаладалада стисаттсас тлапсаслда тсалассатт остилистос стипастдал	480
	GENCTRICACO ATOGRGATOC COGATORGOC CTGCTGGAGC CAGAAGAACA GCCCCAGCCC	120	TCTTGGGAT AAGATCGACA GACCTGAAGC CATCAGTGAA GAGAGGTTGC GAGACATGTT	540
	AGGIGGGAAG GAGGCAGAAA CICGGCAGCC IGIGGIGAITI CIVITIGGGCI GGGGIGGCIG	180	TOSTITATAT GGTCAGACAA CAGGAAAGGG GAGTATATCT CTGAAAGAAC	009
35	CAAGACAAG AACCTTGCCA AGIACAGTGC CATCTACCAC AAAAGGGGCT GCATCGTAAT	240 35	ACCCITIAGAA GITITICANGI GIAQIGIGCT CAAAAGACAA GGITIACOQAG	099
	CCGATACACA GCCCCGTGGC ACATGGTCTT CTTCTCCGAG TCACTGGGTA TCCCTTCACT	300	CHOSANICGOA CAGTACATING ATTRACACAA ACTCACATING GTTTCCAGGIC TCAACGITTCA	720
40	TOSTOTITIG GCCCAGRAGC TGCTCGAGCT GCTCTTGAT TATGAGATTG AGAAGGAGCC	360		0 20
?	CCTGCTCTTC CATGTCTTCA GCAACGGTGG CGTCATGCTG TACCGCTACG TGCTGGAGCT	420		
	CCTGCAGACC COTOSCITCI GCGGCCTGGG TGTGGTGGGC ACCATCTITIG ACAGCGCTCC	480	THE THE PROPERTY OF THE PROPER	
45	TOSTENCAGE AACETGSTAG GSSCTCTGGG GSCCCTGSCA GCCATCCTGG AGGSCCGGGC	540 45		006
	COCCATOCITG COCCEPTING TOCTIONIDG CTTTROCCTIG GIOGRAPIC TOTTICCACOT	009	ACTIOCITY TALASTITY ANTENANT CALCECAGE CATTIGINA AGAICACT	0 66
Ç	CCTGCTTGCT CCCATCACAG CCCTCTTCCA CACCCACTTC TATGACAGGC TACAGGACGC	099	ICCAGCAGIA CATITICANIC ACTITITIANC ANLATAGAMC INTONACCAT ATTIMAMAC.  O DESIGNATION DESIGNATION RESISTANCE RESISTANCE REPORTS ASSETTING A	1020
3	GOCTICTICOC TOGGCCCAAGC TCTACCTCTA YTCAAGOOCT CACGAAGTAG TCCTGOCCAG	720		1000
	AGACATAGAA CGCATGGTGG AGGCACGCCT GGCACGCCGG GTCCTGGCGC GTTCTGTGGA	780	SALEMENTS AND APPLICATION APPLICATION AND APPLICATION AND APPLICATION AND APPLICATION AND APPLICATION AND APPLICATION AND APPLICATION AND APPLICATION APPLICATION APPLICATION AND APPLICATION APPLICATION APPLICATION APPLICATION APPLICATION APPLICATION APPLICATION APPLICATION APPLICATION APPLICATION APPL	0000
55	TITICATOTICA TOTOCACAGO TCAGOCACOT COGTGACTAC COTACITACT ACADAAGOT	840 55		0027
	CHOTOTICAAC TITCATGCGCA ACTGCGTCCG CTGCTGAGGC CATTGCTCCA TCTCAMCTCT	006	TIGGGANAA AAAAAAAA AAAYTGGA	1228
9	OCTCCAGAAA TAAATGCCTG ACAKCTCCCC ACAAAAAAA AAAAAAAAA KCTCGAGGGG	09 096		

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(2) INFORMATION FOR SEQ ID NO: 280:

 $\varepsilon$ SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1327 base pairs

S

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 280

5 TITICIGATIT TRACATGACC TIGAGCAGGT TIGCATATAA TIGAAAGCGA TIGCCCTICIT GOCGOGTOCA GEAGATCOTO GGCGCCCTCC GCAAGGGCGS CGGAGACCGG TTACAGGTGA TOTOGOGIOT COGGACAGGI GAGCACCOIG AIGAAGGCCA COGIOCIGAT GCGGCACOIG

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SEQUENCE DESCRIPTION: SEQ ID NO: 281:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 799 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

20 2 COCTOCTICA COACTATIAC CCAATIGAGA TOGACOCACA COGGACOGIC AAGGAGAAGO TACCTCATAT GGTOGAATGG TOGACCAAAG CGCACAATCT CCTATGTCAG CAGAAGATTC CTTACAATAT TCTOGATAAT AGCAAGATCA TCAGTGAGGA GTGTCGGAAA GAGCTCACAG

> 300 240 180 120

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TAATCCTTGC

ATGTTGGGTT

ATTICCAOCCA AAGACAITTIC AAGTIGCCIGT AACTGATTTIG TACATATTTA CIGCIGAACC CAGCCIGGGC CIGGATOCIC TOTGAATACA TIAICTIGGG

300 240 180 120

COCTITIOSCY GARGECTARY TECREAGREET CITOTITYTY GAGAGAGACT GAGAGAACCA

TOGIATICAC AGCCIGGATG ACCCICACGG CCAGCCCIGG GGIGITICCCC GICACIGIGI TUACCCIOCO TACAGOTOG AGOTUAGATO ACTOCOCOCT CUACOGTUAC TOTOAGUAGO

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25 30 TCOTOTOTAA CTACATOGAT TTTAATGAAG ATOGTTTTCT CCAGGGATTT AAGGGCCAGC TIGGIGATAT CCIGGAAGAA AITAICCGAC AGAIGAAAGI GIICCACCCC AACAICCACA 540 600

TIGAGGGCAA AACCAATIGIC AIRCTIGCIGG GAGACTCIAT CGGGGACCTC ACCATGGCG TEATACACAC ATACAACAAG AACAGCTCTG TGTGTGAGAA CTGTGGTTAC TTCCAGCAAC 720 660

ATGGGGTTCC TGGTGTGCAG AACATTCTCA AAATTGGCTT CCTGAATGAC AAGGTGGAGG 840 780

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GATGTGGTCA ACGGGCTACT GCAGCACATC CTGTGCCNAG AGCGGCGGGA NCGCTAACAT GGACTCCTAT GACATCGTGC TGGAGAAGGA CGAGACTCTG GGGGTCCAGC TGGAGATGCA

AGGCCCCTIGA AGGCGCAGGC TCCNAAGKCC SCTGCAGGCC GTGGTGAGGA GGGGCGCCTC CCCAGAGTCT GETECECET GAACACAGAG CAGAGECAGG GTGGCCAGCA GTGGETGGGT

CCTTCCGCGC CCCTCCGTCC TOCTITICCCT GAGCACCTTC ATCACCAGAG GCTTGAAGGA

ACCCCGCCAT GTGGCAGGGC

GGATTGTCTA

CICCAGGGAT TITCTTCAAA ATTITTAAAC ATGGGAAGTT ACAGGCACTO TICCIOGIGA ACCITIOGACC ACAGCATOTO

CARACARATA TARTOTOTGA AACAGATCAA AATTTTTAAA ATGAAAAAAA AGCTGCTCTG

GGTAGAACCT GGACCTCTTG GCCTGGGGGC ACATGGGATG

TGTGGGTCGG

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1327

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CHOCCHOCA TOGGTCTTGC TTTCCTTTAT ATGACTGTCC TGGGCTTTGA CTGCATCACC

180

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OCTGAGECET TEEGTAEETT CEGAGATOGA TOGOTETEET ACTACAACEA GEETGTOTTI AAAGACTOTA ACATOCATGA GOTTGAACAT GAGCAAGAGO OTACTTOTOG CKSCCAGATG

TCAGCTATAA CTOGAATAAT GOGAACTOTA OCTTTTACTT GGCTACOTCG AAAATGTOGT

300 240

ACAGGOTACG CCTACACTCA GOGACTGAGT GOTTCCATCC TCAGTATTTT GATGGGAGCI

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1200 1140 1080 1020

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€ 0

SEQUENCE DESCRIPTION: SEQ ID NO: 282:

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1260

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SEQUENCE CHARACTERISTICS:

(B) TYPE: nucleic acid (A) LENGTH: 2196 base pairs STRANDEDNESS: double TOPOLOGY: linear

INFORMATION FOR SEQ ID NO: 282:

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INFORMATION FOR SEQ ID NO: 281:

AGACCTICIT CAACACACIC TACCATAACA ACATICCCCT TITCAICTIT TCIGCGGGCA

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AGAAOTTICA GATAGECECAG GIGGITAGAG AGICEAATGE AATGETEAGG GAGGGATATA

420

25

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TIGIOGATIT TATOTOTOTO TOSCITAATA ATCATAGTAA CAACAATAAT ACCITTITCT

CCATTITICT TOCAGGAAAC ATACCTTAAG TITTITITIGT

TITICITITIC

ITTITTTTGIT

ABABARTABA ATAGTCACAT TTTTBATACY ABBARATGGA

799 780 720 660

35

CHANAMAGT CGAGGGGG TITIOTITIC CITTATGAAG ACATTIGAAA CAGICIGCAC CITIGATACG GIATIGCATI TCCAAAGCCA CCAAICCATT tigeoteset gatossstig aastitostt tostietist tieaseeeaa tatstagaga

AGAGAAGGAT TOOTOGATOT AGOTOGTOAC GAOGATOTTT TOACCAAGOT CACAGGAGCA 600

480

GCCCTTCAAC CCCACCTTGG ACTTGAGGAC CTACCTGATG GGACGTTTCC ACGTGTCTCT

TAAAAATCTA TICAGAAATT GOTCCAATAA TOCACGIGCT TIGCCCIGGG TACAGCCAGA 420 360

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120 180 240 300 360 420 480 240 900 099

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	209		310
	TTOSTTCCCA CAGGICTICAT CTCAGGATTG GCACAGCTTT CCTCTTTGAT CTTGTGTGTG.	360	TAAICCITIG CITCAICITI CIACAGIAIG ACAIAAIGAT TIGCIAIGIT GIAAAAAICTI
	ATCTCTGTAT TCATGCCTGG AACTGCTG GACTTGTCCG TTTCTCCTTT TGAAGATATC	420	tgtaaaaat ttctatataa aatatttgaa acttaa
<b>.</b>	CGATCAAGOT TCATTCAAGG AGAGTCAATT ACACCTACCA AGATACCTGA AATTACAACT	480	\$
	GAMAININGA TOTOTANING STOTANITOT SCIMINITIG TOCOGGAGAC ANSTOCTIGAN	540	(2) THEODIMATION FOR ERO TO MO: 281.
2	TCTGTGCCCA TAATCTCTGT CAGTCTGCTG TTTGCAGGCG TCATTGCTGC TAGAATCGGT	. 009	(i) crottman characteristics:
2	CITIGGECCI ITGAITITAAC TGEGACACAG ITGCTGCAAG AAAATGTAAI TGAATCTGAA	. 099	
	AGAGGCATTA TAAATGGTGT ACAGAACTCC ATGAACTATC TTCTTGATCT TCTGCATTTC	720	
15	ATCATGGTCA TECTGGCTCC ANATECTICAN GCTTTTGGCT TGCTCGTATT GATTTCAGTC	780	
	TCCTTTOTGG CAATGGGCCA CATTATGTAT TTCCGATTTG CCCAAAATAC TCTGGGAAAC	840	GENERAL GENERAL GENERAL CONTRACTOR GENERAL GENERAL CTGGNGCTCC
۶	AACTETITG CITGCOOTEC TGATGCAAA GAAGTTAGGA AGGAAAATCA AGCAAATACA	006	30 APPRING GANGEAGER CAGAMYCOAT CITICAATITIC TGCTAAAATA CITICIDACIC
3	TCTGTTGTTT GAGACAGTTT AACTGTTGCT ATCCTGTTAC TAGATTATAT AGACCACATG	096	
	TOCTTAITTT GIACTOCAGA AFTICCAATAA ATGOCTGOGT GITTTTGCTCT GITTTTTACCA	1020	ATARIGARIC TCHARLANGE TCTOTATIC ATCTACC CATACONS CONTRACTOR
25	CAGCIOTGC TIGAGAACTA AAAGCIGITT AGGAAACCTA AGTCAGGAGA AATTAACTGA	1080 2	GIRARIUITA AGCIGGACCI TOGCACCCI CITACCATUA AGCACCITA CITACCATUA CITACCATUA CITACCATUA CITACCATUA CATACATUA
	THARITICCC THATGITIGAG GCATGGAAA AAAATTGGAA AAGAAAAACT CAGTTTAAAT	1140	AGACICCCC MITRIGGGA ACAMANITC MITTATICE CIRTITATION ANGESTITES
	ACGGAGACTA TAATGATAAC ACTGAATTCC CCTATTICTC ATGAGTAGAT ACAATCTTAC	1200	
93	GTAAAAGAGT GOTTAGTCAC GTGAAFTCAG TFATCATTTG ACACATTCTT ATCTGTACTA		30 CIATCACTOT TOGCATOCTI TOGTOACTOG GAGATOCTIT GGGGGTGGG AGGTOCTION
	THE THE PARTY AND TAXABLE AND	• • • • • • • • • • • • • • • • • • •	GTCCCAGGCT AAAGGAAAAG CTTCACAAGG GTAAGAGCCA CAGAACCCTC GGCAAGAAAG
3	GARTICAGAT ATGICAGTIT TCIGCAAAAC TCACILIGI TCAAGACTAG CIAATITATI		GCCGCTCAGG GAGAATGAAT GGTACAGAGA GGAAAGGAAG GAAAGGGGGT GGAACAGAGG
33	TTTTTGCATC TTAGTTATTT TTAAAACAA ATTCTTCAAG TATGAAGACT AAATTTTGAT	1380	3.3 TAGAAGGCAA GGAAGGGATG CCGCACTGGA GACCGATGGG GACACTCTAA TTGTGCAAGA
	AACTAATATT ATCCTTATTG ATCCTATTGA TCTTAAGGTA TTTACATGTA TGTGGAAAAA	1440	GGGAGGANCT TOCTTCTTGA ATGCTGAACA CAGCTAGTCT GAACCTTCCT TGGAAAGTCC
6	CANANCACIT ANCHAGANTT CTCTIANTANG GTTTATGGTT TAGCTTTANG AGCACCTTTG	1500	40 местетттес ссатесатые воссыястет ссстесамые смесымитет весттетите
•	TATTITIAIT ATCAGATGGG GCAACATAIT GTATGAAGCA TATGTAGCAC TTCACAGCAT	1560	AGGAAGGAAA AGTATCCATC AGTGTGACAA GAGGTCACCT TCGAACTTGC ATGAACTC
	GOTTATICATO TARGCTICCAG GTAGAAGCAA AGCTOTAAAG TAGATTTATIC ACACAATIGAC	1620	TESTS AND ANAMARISES TOSTAGAAGT GAGGAATCGCC TAGTICTTAGG GCTGTCCGTT
45	TECATACAGA CITCAMATAT GICAATAGIT TOGICATAGA ACCTAGAAGC CAMAAGCCAC	1680 4	45 managages constitutes appointments a parabaches apprendence
	ACAGAAGGGC AAGAATCCCA ATTTAACTCA TGTTATCATC ATTAGTGATC TGTGTTGTAG	1740	CANCIDICA TANGARGE TIXCASCOTA GITCATCACC CAGACATGA
ç	AACATGAGGG TGTAAGCCTT CAGCCTGGCA AGTTACATGT AGAAAGCCCA CACTTGTGAA	7800	S) cosecuent cacestabae macacatric meacantam contemporation
3	GOTTITICITI TACAAMICAC ITGAITITAAC ACACICAGGI AGAATATITI TAITITIACI	1860	
	GITITATACC CAGAAGITAT TICIACATIG TICIACAGCA AGAATATICA TAAAAGIAIC	1920	CHARLESTON VICENCE STREETS PROBLESS PRO
55	CCTTTCAAAT GCCTTTGAGA AGAATAGAAG AAAAAAAGTT TGTATATATT TTAAAAAATT	1980	55 mmchannac acamemen controllera carabacare recta
	OTITIANANG TCHOTTISCA ACAIGICIST ACCAAGATOS TACTITISCST TAACCOTTIA	2040	
9	TATOCACTIT CATGGAGACT GCAATAGGTT GCTATGAGCA CTTTCTTTAT CCTTGGAGTT	2100	09

720

780 840 900 960 1020 1080 1140 1185

(2) INFORMATION FOR SEQ ID NO: 284:

(C) STRANDEDNESS: double (B) TYPE: nucleic acid

S

20 15 5 AAGCTTGGTT TGTACAAAAG CAAGGTGGGA GTCTATTTTT GTACATGAGA TACATCACAC CCACCACAAA TIGIGIACAT AGTCITCAGA TGATACCACC AAGAGCTGGT TTACCTGTGG AGCAGCCTGG TGCAGTGCCC TGTCATCAAG ACAAACCCAC GGTCCTMCTG ACCCAAAGOG GAGGOTAGCG GGAGGGTAGC AGGTGAGTTC CTAGGGCTGG TCTAGGCCTG TGTTATATGT CATATTTAGC OCCAGNATIO TGAAGTGAGT CIGAGTIGIT TACACIGATG CCTTCCCTGC STITTITATAT ATGACCTITG CCTTTCCCCA GCTCCCAACC GGTGCCTACC

300 240

ATTICICITIC TITIGIATITI AGCACAGIGT ATGCACCITC ATTTAAATAC ATCTGTGTGC 360 420

GTTCAGCTGA ATACAGATAC AGCAGATGGA GTCCTGCAGC CCAGGAGACA CCCTGCATCC CTGCTAATAG 540 480

TGTTTGCCAC AAGTATTAGT GAGTCTTCCT TATTAATATT TICATTTCAG AAGACTGAAG 660 600

780 720

840

900

CICGITICCCA AIGATAGAIC ACICCIGITG ACCIGGIATG ICIGCITIGCI IGCIGCITIT 960

CCTTGCTTTC TCTTGGAAGA GGAAAGGACT CTGGTCAGGC CCAGGCTGAG TGAGATGAGC TOCAGOTOGO TOATGGOOTT OTTAGAGOAG AGAGAGGAGT ATGTOATTIT ACTAAGITOO

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THERECARGA TOTCHOTAGE ATCHCCTCCA CCCTCCTTGG ACATGAATCC TCCATGGAGG

TGAACATGCT GAATCATCTC CAACAAAACC

TOGACTOTOT GATCCCAGCT CTOGAGCAAG CTGTAGACGG ACAGCAGGAC ATTGGACCTC

GAGGACITTO CIGIOCIGOG IGAAGGCCIG GIGCAGACIG IGGAGGCCCG AAGIGAGAGG COCATOGIGA TGTCTCGAGG TGGGGAGCTA GTAATGGTGC CCAATGTTGA AGCAATCCTG GICCITCIGGA AGCCCACCAG CTITCCAGCCG CAGCCCTICA AGGCCTCCAG CACAAAGGGG GAGGCCTTTG GGGATCTGGC CCTTTTCTTC TATGACCAGC ATGGTGGAGA GGTGATTGGT TCATCCCTGA TGCCCGTGCT GOGTNATGAT CCTNCTCAGC TCTATCTGAC GCAGCTCAGG AGGETTOTOG ACTOGECAGY TOCCTOCTTO TOCCOGGGGC TOCTCAGGCA GEOGGGGCCC

> 660 600 540

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CCCTIGGACA TITACGACGT GCTGATTCGC CTGTYTCCTC GCCATATCCC GCGGCACCGC

480

420 360 300

ITRAGRARGE RECICATIGA TECEEGOGA CETOGOGRER TERGGRERGT CITTEEGOCEG GUACCOTOAG COCAGATOOT GOAGCAGOTT GTOGTOOTOG CAGCTGAAGO COTGCCCATG CICHCCGIYA IGGICATIGI TACCCCCCAA GACCGCAAAA ACICIGIGIG GACACAGGAI CAACCCCCTC TITIGTCAACC TCAATAATGA GCTCACTGTG GAGGAGCAGC TCGGGCACAG

240

180 120

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TICCCCCCAG GITIGGCITICC TICGATICCT TITCTIGGIA TCAACGITIG ATTIGGAAGAA

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SEQUENCE DESCRIPTION: SEQ ID NO: 285:

TTATIGACTIC CICAAGAATA IGTAGCIGCT AGGGGTAAA ICAAGGCATC ACAATITCIG TARACARACA TITATIGUAGO CARCACICCI TIGUAGATICCA GARACIGAGO CACARIAGGO

> 1140 1080

1200

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ATTOGGGCAG

GGCATGGTG GCCCATGTAG

CAGAAGAGGA GTGGGAGCCA GCTCAGAGAA GGAACTGAAC

CATCCACCTA CTCACCATCC TITCICICIG

960

900 840 780 720

COCTGCGATT TOOCACTOOT

TICAGEOGGE AGGAATAGGE IGIGAATIGE TAGEACTITT TITITITAAG CAATTACTIT

TIGACTIGIT CCICIGAAAG IGCAAGAGGC GIACACCIIT CCCAAAIGIA GACTAGAAIC

IOCAGGATOC CACCCACTOT ATAGITICIGC TITICCCAGAG AGGAAGAACT TITAGAAACC

AAATGATCTT 'AATTGTTATT GCCCACCCCT GGCTTTTCCG GGTAGAAAAT TCACAGTAGG

8

ACATATCTGC AACCAGAGCA GCCACCAAGC ATTACTTAGC AGCAGGAAAA TGATTGTATT AATGATTOTT AAGAGAGAGT GCTTGGAACC ATGGGTTAAC AGGAAAGGCT ACCTAACTTC

> 1500 1440 1380 1320 1260

8

TOTOGOCCOTT CTGAGTCACA TCACCGACAC TGAGCAGTGG AAAGGGGCTA TATGTGTATG

AGCATGCCCT

TCACTGCAGT GTCAGGCCTT TAGATGGGAC

CCAGCGAAAA

1260 1200 1140 080 1020

GGATGTGTGC GCCTGGACCT

CTGGGAGAGG

CICICICICC

TICCIGCIGC TICTTAGICT CCAGGAGATC Tercerosce CAGCCCCAAC

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TOGITIONOC CATTIOGGGA TAAGGIOCIG AAGCCAGAGC ATTIOCAGIT IGITIGAGGC

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TOGTACTOTG TCTOTOCCTO TOTOTOTOTO TGATAGTCAC TCTTGCATGG CTTCCATGTC

TGIGITCCTC

CAAAGCTGAT

ARGCIGGACT TIGGGGCCAT CICIGGAGIA TIAGCCCCCT TITTIGCTIGG

ACICITICCT CITICITICS CACCIAACIG ACCICTICG GAIGACITCC

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GCATATATGT GIGIGGGIAT GCATATATCT CICATCIGTA GITTCCAAGA

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180 120 6

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1795 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

INFORMATION FOR SEQ ID NO: 285:

KITICACCGG GGIC

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: (D) TOPOLOGY: linear

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 1634 base pairs

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WO 98/39448

TGAGITICCIG IGIGICCAAA ACIGAGGCAC CATGITICITT GAAAACATGC CACCICAAGG

CTOGOCOCGO TOGOTUADAC CTOTAATCCC AGCAYTTTOG GGAGGCCSAG GOCOOGGCOG

1620 1634

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	513		514	·**
	ABMACAPTAL APPCABACEAC PACEARTOPP PROPERTIES MECHANISM PRESIDENT	. 0011	TTAAAACCIT ATAAACTA	828
			-	
ν.	Gacagtogaa aagaaccgag gacaggaaag gattgggtag gtgaaggggt caggggactg	1380 5		
•	GTAGTCACCC ANTCTTGGAG AGGTGCAAAA AGCACTGGGG GCTACCCGTT AGCTGCATCT	1440	(2) INFORMATION FOR SEQ ID NO: 287:	·:
	GCCCTGGCTG TTTGCCCGTT CATGTCACAA ACTGCCACTA CTATGTACCT GCAGTGGGGT	1500	(i) SEQUENCE CHARACTERISTICS:	
01	TECHGAGATG GOGGAGACTE AMOTETTACT COCCAGGAGC TECCAGGGCC CAAGGAGGAG	1560 10	(e) (c)	
	AATGCTGCCT CCTTTCAGTC 1637CTAGAC CCACTTTCTG GTAGCCTCTC 19CTTCCTGT		(D) TOPOLGGY: Tinear	-
9	ANTICTOSCY OTTITICCAG ACTCAGCTCA ANTIGETOCC CTCCTTAGG CCATCCCTCG	1680	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 287:	
2	CCCCCAGCCT GAGGIGATCT TTCCCTCCTC TGAACTATTA GAGCAGTTAC TGTCTGTTCA	1740	GAATHOGGCA CGAGCGCGC CATGCCGCTC CTGCTTTCGG TGCTGCGTGT ACTGCTGGC	09
	OTTOOTITOS CAGGCACACA CAGTGGCATA AATTCTATTG TITTGAACTC TGATT	1795	GOCTITETICS COCTOSTISS STRESCOAAG CTCTCGGAGG AGAICTCGGC TCCAGTTTCG	120
70		20	GACCOGARGA ATGCCCTGTT COTGCAGTTT GCTGAGGTGT TCCCGCTGAA GGTATTTGGC	180
			TACCAGCCAG ATCCCCTGAA CTACCAAATA GCTGTGGGCT TTCTGGAACT GCTGGCTGGG	240
	(2) INFORMATION FOR SEQ ID NO: 286:		TIGCIGCTGG TCATGGGCC ACCGATGCTG CAAGAGATCA GTAACTTGTT CTTGATTCTG	300
25	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 858 base pairs	25		. 360
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double		CCAGCCAITS TCTGCCTGGG GTTCCTGCTG CTGCTGAATG TCGGCCAGCT CTTAGCCCAG	420
30	(D) TOPOLOGY: linear	30	ACTIANGAAGG TOGTCAGACC CACTIAGGAAG AAGACTCTAA GTACATTCAA GGAATCCTGG	. 084
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 286:		AAGTAGAGCA TCTCTGTCTC TTTATGCCAT GCAGCTGTCA CAGCAGGAAC ATGGTAGAAC	540
	TCTGCTTTCG GTGCTGGGTG TACTGCTGGG GGGTTCTTC GGGCTCGTGG GGTTGGCCAA	09	ACAGAGICTA TCATCTIGIT ACCAGIATAA TATCCAGGGT CAGCCAGIGT TGAAAGAGAC	009
35	GCTCTCGGAG GAGATCTCGG CTCCAGTTTC GGAGCGATG AATGCCCTGT TCGTGCAGTT	120 35		099
	TOCTIGNOSTIG TTCCCGCTGA AGGTATTITGG CTACCAGCCA GATCCCCTGA ACTACCAAAT	180	AUSTIANGCA TAITIAACATT CCTCATGTCA TATGAMAATA CAAAATAACC AGAAAAGAAA	720
40	AGCIGIGGS TITICIGGIAC TGCTGGCTG GTIGCTGCTG GTCATGGSC CACGGATGCT	240 40	TTTAAATCAA	
•	GCANGAGATC AGTAACTIGT TCTTGATTCT GCTCATGATG GGGGCTATCT TCACCTTGGC	300		840
	AGCICTIGADA GAGICACTAA GCACCIGIAT CCCAGCCATT GICTGCCTGG GGITCCTGCT	360	APARABABABAB MARABABABABABABABABABABABABABABABABABABA	006
45	GCTGCTGAAT GTCGGCCAGC TCTTAGCCCA GACTAAGAAG GTGGTCAGAC CCACTAGGAA	420 45		
	GAAGACTCTA AGTACATTCA AGGAATCCTG GAAGTAGAGC ATCTCTGTCT CTTTATGCCA	480	AUTOMOTOR CITCUIT	
Ş	TOCHOCTOTC ACAGCAGGAA CATGGTAGAA CACAGAGTCT ATCATCTTGT TACCAGTATA	540		
3	ATATICCAGOS TCAGOCAGTO TTGAAAGAGA CATTITOTICT ACCTOSCACT GCTTTCTCTT	009	(2) INFORMATICM FOR SEQ ID NO: 288:	
	TITAGCITTA CIACICITIT GIGAGGAGIA CATGITATIGC AIAITAACAI TOCICAIGIC	099	(1) SEQUENCE CHARACTERISTICS: (1) TEMOTH: 1517 base pairs	
55	atatgaaaat acaaaataag cagaaagaa atttaaatca accaaaattc tgatgcccca	720 55	( a (	
	AATAACCACT TITIVATGCCT TGGTGTAAGT ATACCTCTGA ACTITITITCT GTGCCTTTAA	780	(D) TOPOLOGY: linear	
99	ACAGATATAT ATTTTTTT AATGAAATA AAACCATATA TCCTATTTTA TTTCCTCCTT	840 60	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 288:	

(A) LENGTH: 3865 base pairs (B) TYPE: nucleic acid

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Ξ SEQUENCE CHARACTERISTICS: 2 INFORMATION FOR SEQ ID NO: 289:

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50 ТТССАВАВАВ ВАВВАВА GGGAGGTCAC CTITGGGIGT GIGIGGGIGI GIGIGIGIGI GIGIGIGGGI GGGACGGGIG GCGGTGTGCC CCCAGGACCT GTAAGTAATA AAATCTTTAT 1500 1440

3 CCAGTTAAAA TCTCCTCAAA ATGTTTGGAT ACCGCCCATT GGCCCCTCAC AGCCACGAGC GTGACCCCCT CTCAGGCACA GCATGACCTC CTGAAGTCGA GCCTGCTTGC TTTGAACCTA

3

GTTCCCCCCA CAGCCAGTG GTCTCCCCCC ACATTTAGAA GCGAAAATGT OTTICATITA AAGAIGITAA TIAAATGAII GAAACIIGGC IGIGGCIACI

ITICCCICCI GCTTCTTAAT

> 1200 1140 1080

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OTOTOTOTO CATOTTOTAL COCCCACCCC CCATTACOGG TAAAGGRAAC CCCAGACTAG GCACACHTIG GACCCAAGTA TOGGCCICTT CTGCCTAGTA CTGCCAAAGG GACTGTTAAG

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AAATTCCAGG AGCCCTGGGCAGCCCTGG NCCCCAGTGC CAAGCCTCAG AGTAAGCAGA

1680 1620 1560 1500 1440 1380 1320 1260

GAAGTACCCC ACCCTCTATG AAGACAGAAT CACTCTCTGC

TICAGACATT TIATITIGAAT TIATGACAGT GATGGGGATT TGACTGAGAT AGCAGCTGCC TCACATTGTG TTCTCTCCTG AGATGGTCCA GCTCACATCC

AACAACACGC AGCTGATTTA GGGAGTGTCC CAGCCTAGCT GGATCAAGGG

8

TAAAANGITA GAAGIATAIA TATACATATA TATATITICIT TAAAITITIIG AGICTIIGAT

GTAAATGAGC TTGACCTAGA GTAAATGGAG AGACCAAAAG CCTCTGATTT TTAATTTCCA GCACAGAGGA CAAAGGAGAA GGGAGGGTCT AGAAGAGGCA GCCCTTCTTT GTCCTCTGGG TECANCATTA CIGGANCIET ATCEIGITAG GAICTICIGA GETIGITICE CIGCIGGGIG AGENACCETG GENATIGGETG GAGGTISGENG AGNACCTIGNE TICTICTITICE CICTICCETICS

1020

960 900 840 780 720 660 600 540 480 420 360 300 240 180 120

ATGTCTAAAA ATCCATTCCC TCTGCCCTGA AGCCTGAGTG AGACACATGA AGAAAACTGT

GACAGOCTAC TIGACOGOCOC CGCIGGCOCO CACATIOCAC TGAACIGIGC GGAIGCCACA 1260 1320

GATATETCCG TTATGCAGCC GCCTCCGGGG GACCACCTCC CTCCCTTTGA GTCAGCCACA 1200

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GTCACCGTCA GCAGTTTOCA GTTTTCCACC TCCMCCCAGT TCCTCCGTGT GGTTGACCCA 1140

AGTITOCCTIG GGATICATIT THATIGIHAGC INGACITIGI CATGCCHGAA ACAAGGCING

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GOGITTETICE CATGGACAAA TIGCCITECA AMAMIGAGGA CATCATGAAG CAAAAACAGA

GOGCCCTCCC GCCGCRGYGA CCTCCAGAGC CTGGGYTANT GCATGCTGAA GTGGYTCTAM

GEAGGAGECY TEAGGAGGG GACCITGAGI TICATINGCA TGGACCIGCA CAAGGGAIGC

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ATCTACCCCA CCCCCTAGGA TCCAGGACTG GGTCAAAGCT GCATGAAACC AGGCCCTGGC

CTGCCCACTG AGTTGGGGAA AGAGGATAAT CAGTGAGCAC TGTTCTGCTC AGAGCTCCTG AMPCANTATT CTTCCCTTGC CTGTGGGCAG TNGGAGAGTG CTGCTGGGTG TACGCTGCAG CICHOCCCI CICCAGGGIG TITTICCACIA GICACIACIG ICTICICCII GIAGCIAAIC TRACCTECCE GEOCETEAC CAAGGETGGG AACAGAGGG ATSTGGTGAG AGCCAGGTTC

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GOSCITGAGI CIGGCAAGGA ACCITGCIIT TAGCITCACC ACCAAGGAGA GAGGITGACA

GOGIGACISC CCCCCAGGGC CACCGCITCT TICTIGATICC ICTITCCITA ACAGIGACIT AGCCTOTOCC ACCTOTOTAG GCAAGCTGGC TICCCCATIG GCCCCTOTOG GTCCACAGO

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TOSTITCCACT GGGGGCAGCT AACGCTGAGT GACAAGGATG GGAAGCCACA GGTGCATTTT

TRACCITICT CICICCICAT TICOGISCAT GICCTITICIG CAGCISCCIT ICAGCACAGS TOTAGOGOGG GAGCTICCTT GAGCAGTOGG CCCAGGCCTG GCCCTCCACA CTTCATTCTC

GICCCCTAGG TATCAGCCIC ICTIACIGIA CICICCGGGA AIGITAACCI TICTATITIC ACTUARGICT TUTUTAGICA ATGRIGOGICA CUCAGIOCIT CIRCOGURGO CIGOGIOGIO

OCTATOSCIT COCNTICCOC TATTOCCCAA GIGÓCAAACA COIGOCCIAC

ATGTGACAGC TGAAAATATC TITOTXGATC CAGAGGACCA GAGTCAGGTG ACTITIGGCAG 840

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TOOCCIOCCO OCTOCIOGAT OCCCIOGAGI TCCICCATGA GAAIGAGIAI GIICAIOGAA 780 720

TCACTOGGCC CTGGATGTCA GCCCAAAGCA TOTGCTGTGC AGAGAGGTCT GTGCTGCAGG TITICOGICIT CACCAGGACA AATACAGGIT CITOGICITA CCCAGCCIGG GGAGGAGCCI

20

AGTCAACAAG TOGAAGAAGC TGTACTCGAC CCCACTOCTG GCCATCCCTA CCTGCATGGG

CAAGGATGGG CGCTTGTTCA ATGAGCAGAA CTTCTTCCAG CGGGCCGCCA AGCCTCTGCA

15

CTCCACCCTC ACCTGTGACT CAGGACCACA GAAGCAAAAG TTCTCACTCA AACTGGATGC GAAGCTGAAG TCCTTCCAGA CCAGGGACAA CCAGGGCATT CTCTATGAAG CTGCACCCAC

CCACCTCACT TGAAGCTTTG CCCACAGGAC AGTGCTGACA GACAAGAGTG GGCGACAGTG

5

CCAGCTOTAG CCCTCAGAAG ACCAGGCAGA GCCCTCAGAC GCTGAAGCOG AGCCGAGTGA

AMAGCAGOCC TCAGAAGACC AGGAAGAGCC CTCAGGTGAC CAGGGGTAGC CCTCAGAAGA

S

CCTTOTOGICA ACTAGTOGGT CCCCCGGGCT GCAGNAATTC GGGCAGTGGT TCTGNOTCTG AAGATACTOT GAGTTOCTOT GAGAGATOCA AAGGOTOCGG GAGCAGACCC CCAACCCCCA 120

<u>X</u> SEQUENCE DESCRIPTION: SEQ ID NO: 289 (C) STRANDEDNESS: double (D) TOPOLOGY: linear

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	CATTOGGAAA GITGCCAACC ACITGGTAGA CCACTAGGIT CICTGTITIC CCITCCCTTT	1740	CATAGTAACT GACTCGGAGT TCTAGAAGTT CCCATGGGGC TTTCATGTTA TCCAGGTATG
v	CCTTITICAAA TCCCACAGITI TCCTGITIGGG GAGAAGCIGT AATTAGCCTA GICCAGGIAC	1800	CGTGGAGGTC CGCAAACAGA ATTGTTACCT CACACCCTGC CTTTAAGAAG TCTGCAATCT
ר	CAGATCCCAG CTAGGGGGGC AGCTGNCTTG GATAACTCCA AGAAAACCTG GGCACCAGTA	1860	TTGACATGGG CACAAAGTAA GCCACATGTG GTTTGCCCGT GGTTGCCGTT CCCCAGTAAA
	TTTTCCAAT TATAAGGACT GTGGCATAAA TTTTTAAATG AGTTATATTG AAACCAGATT	1920	TITIAAGITC COGCICCITIC AGIATCICCT TCAGCITICIC ITCCCCCAGA ACCICCIGGA
9	TETECRACITS CCAAGGGAAG AAGGTAGGG TGGACTCCCT GCTGTGGCC AGCCCTTGTT	1980 10	GOTTICCOGGT GATAAGGTGC AGTITICTCTT CAGGGCTGGG AGGGTCCCCC ATGGTCCGCT
	AGGGGTTGGT CTCTCACTGC AGCCAGACAG GATGATCCTG GGTTCT0GGG AGGGTAAGCT	2040	ACCCCTOCTT COCCCGCTCA GCCCGGCACC AGAGCCCCTT CCTGGGTCAC CGTCGCCGGC
<u> </u>	GCCCCTTGCC GAGTTCTGCA CCGAATAAAG AGTCCAAACC CGCTGCTTCC GTGTCCTGAG	2100	GCGTGCCGGG AACTGTCACG CGAGT
2	AGATOGGTAA ATOGGTGATG GATGGAGCAG ACTGAAGAGA CAGCAGATGA CTCAGTOGTG	2160	
	GAAGAAGGG GGAAGATGCT GGCTGGCTA GCTAATGTTC CCCCCTTTCA GGGATTTACA	2220	
20	GGAAATGGAG CCCAGCTTGG TCATGAAGTT GOTTTGCTTC CACTGTGCGA TGCACTCCTC	2280 20	
	AGANATITIS ANGICAGOCT GONACTICIC GANGACITIC ITCTIGGGCT IGAGOTOCIC	2340	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1910 base pairs. (A) TATOR: ALPHANISTERISTICS AND AND AND AND AND AND AND AND AND AND
75	ATCHGGITGG CCCTTTTCAT AGCCCTTCAC AAACAGGIGC TCACCAGGAG CAGAGCCTGC	2400	
3 .	COGRAGOTIC AGROSTICAA CTGGCGGTTT ATCCCTTCTA TAGAAGCACA CAGAAGCATG	2460	
	CCTTGGGACT CGACTCCTCT CATCTTCTGG GGTTTCAGGT TGCACAGCAC CACTACCAGC	2520	(XI) SEQUENCE DESCRIPTION: SEQ ID NO: 230:
30	CTETICETECA GITICETECT GOGGAGAAC TGTACCAGGC COCTCACCAC AGTCCGTGGT	2580 30	AGGENGAGGA GGAGAGGGG TCTGCOCCG GCCGCTACCC AGAGCCAGC GGACGCAAC
	TCAGCTITCCC CCAGTICIAT CITICICIACA TACAGGCIGT CTGCATCTIGG GTGCTITCTCC	2640	ACGGAGTIGGG CTGTCCCCCGA GCCCAGCCCC GAGGGAGCCC CCCCCCCCCC
ý	ACHGIBATGA TETTICCCCAC ACGATATICC ACCCGGANTO GANGACCTC CTCTGGTTCT	2700	GOGCTIYCCA GCCAGCCCGA CTCCTAGGAG GAGGGGAGGC GGGAAAGCAG CTCAAGCCTC
c C	GANTICITIOS CAGOCCITIOS OCCATIGOCT TCTGCTITICA GOGATICTIOSO TAGOCAGOC	35 2760	ACCCACCACC CTGCCCCCAG CCCCACACT CCCAGGCTCC TCGGGACTCG GCGGGTCCTC
	TOGCCAOTIT TITCAGGGCA GGGGTATIAA ACTITICCCG GAITGGAICC AGCAACTIGE	2820	CTGGGAGTCT CGGAGGGAC CGNCTGTGCA GACGCCATGG AGTTGGTGCT GGTCTTCCTC
6	TCAGTECEAC TTCAACAGAA TTCTTCAGGT CTCCAGGATG TACAACCTCA GCAGCAAAGT	2880 40	TOCAGOCTEC TOSCOCCOAT GOTCOTOSCO AGTOCAGOTE ANANGOAGA GGAAATGGAC
	COTTITICAG GICCAGGIAA GCIGISTAAG TITITGITICC ACCCATTIC TCANCIOGTA	2940	CCFFFFCATF ARGATTACCA GACCCTGAGG AFFGGGGGGAC TGGTGTTCGC TGTGGTCCTC
	GGATCACAAA CTCGGAACTTA AGGGGAAAAA GGACATGCTT GATGAAGGAC AGAACCCCAT		TICTCGGTTG GGAICCTCCT TATCCTAAGT CGCAGGTGCA AGTGCAGTTT CAATCAGAAG
5	TOTTCTCCAC ATTICCTGGC TCACAGAAGG CCTTCTTCAG TITTITCTTC ACATCCTCT	3060	CCCCGGGCCC CAGGAGATGA GGAAGCCCAG GTGGAGAACC TCATCACCGC CAATGCAACA
	TOCGATICAAG GAGATICAATO TITGACTICOT COTICIGAAGA GOTICATITITG CIGOOTIGITIA	3120	GAGCCCCAGA AAGCAGAGA CTGAAGTGCA GCCATCAGGT GGAAGCCTCT GGAACCTGAG
50	ATCCTGGAAC CATAGATTC ATCAGATGGA CCCGTTTTGA ATAGCCAAGT GCAGGGAGGT	3180 50	GCGGCTGCTT GAACCTTTGG ATGCAAATGT CGATGCTTAA GAAAACCGGC CACTTCAGGA
	ACTICICISC NAGGIGNA ATCITICICT GATCHATIGC TCCAAATIGG GCATCHACTE	3240	ACAGCCCTTT CCCCAGGAGA AGCCAAGAAC TTOTGTGTCC CCCACCCTAT CCCCTCTAAC
ų	THAMPACT TECHTOCAM GOTTGCAGTC COCOGRAPA GAGGOCACTC AGGAAAGGGT	3300	ACCATTOCTIC CACCTGATGA TOCAACTGAC ACTTGCCTICC CCACTGCAGC CTGCGGTTCCT
S	SCINCERCCIG CITTACCACC TEACTICEAG CCITICITISSA ATCORDOTOR GIBACCACOS	3360	OCCCACCTCC CETCANDIST STOTETISTS GIVEN GACTOTISTS GITTISCTAL
	AGGIGAGTET GTACACATET AGTGTGTACT CTTTGCTGAG CTGGTAATCA GTGCCTTTGA	3420	TOTOCTOTT CTGCCTACTT CTTTCTGCAT GSTATTCTGT TTGTTAGTGA ACTGTGGACT
9	TRANCITIONG CITCTCCANG GGCACACCAA TGCTCTCCAG CATTGCTTTG ATCACATTCT	3480 60	COCTITICCEA GOCAGGGGCT GACCACATG GCCATCTGCT CCTCCCTGCC CCGGTGGCCC

TOCATOACOT TOTOCTOCTA GUAGGOTOCT TOTTOCCODA GACCAGOCCO CTOCCCTGAT

1020

20 30 25 5 5 S CCAATAACTC CATGGGCTCT GGGACCCTAC CCCTTCCAAC CTTCCCTGCT TCTGAGACTT GOTTTGCAGC ACTITISTICAT CATICITICAT GGACICCTIT CACICCTITA ACAAAAACCT AGGCTCCGTG CAGCCCTTGG GAACAGTGAG AGGTTGAAGG TCATAACGAG AGTGGGAACT TOCTTCCTTA TCCCACCTGA TCCCAGTCTG AAGGTCTCTT AGCAACTGGA GATACAAAGC TTAGGGATGC GTAGGGTAAG AGCACGGGCA GTGGTCTTCA GTCGTCTTGG GACCTGGGAA CTOTGACCCA TIGCIGITCT CIGITATCGIG ATCIATCCIC AACAACAACA GAAAAAAGGA CATCCTTTGC GOCAATAGIT GAAGGACICC TOTICCGITG GGGCCAGCAC ACCGGGATGG ATGGAGGGAG ATTICCAGGCC TTCCCAGGGG AAGGAGCTGG ATAAAATATC CTTTGTTTCM TAAAAAAAA AAAAAAAAAAA AGGGGGGG CAACCCAGAT CCCGCCCCCC CTGTCCTCTG TGTTCCCGCG GAAACCAACC AAACCGTGCG AGCAGAGGCC TTTGCTTCTC TGCCTACGTC CCCTTAGATG GGCAGCAGAG GCAACTCCCG CAATCTACAG TCTGCCTGTC GGTGGTCAGA GCGGTGAGCG AGGTGGGTTG GAGACTCAGC CCCAGCTCAT CCAGATGCAG ACTACAGTCC CTGCAATTGG GTCTCTGGCA CAGGGCTTCT ACTOTOCCCC TGGGGAATGT GICCCCTGCA TATCTTCTCA CITICOACGAG GAGIICCCCAT CIGCCCCGCC CCITICACAGA GCGCCCGGGG TOAGCCCAGC GITGACGTCA GGCAGGCTAT GCCCTTCCGT GGTTAATTTC 1860 1800 1740 1680 1620 1560 1500 1440 1380 1320 1260 1200 1140 1080 1910

23

AGCAGAGGGC ACATOTGATT GTTATOGCAG CAACCAACAG ACCCAACAGC ATTGACCCAG CTCATOGCGA GGTGGAGCGG CGCATTGTAT CACAGTTGTT GACCCTCATG GATGGCCTAA ATGCTCCTGC CATCATCTTC ATTGATGAGC TAGATGCCAT CGCTCCCAAA AGAGAGAAAA GCAAATTGGC TGGTGAGTCT GAGAGCAACC TTCGTAAAGC CTTTGAGGAG GCTGAGAAGA 20

GAGCTOTAGC ARATGAGACT GGAGCCTTCT TCTTCTTGAT CAATGGTCCT GAGATCATG

AGCCTCCTAG AGGAATCCTG CTTTACGGAC CTCCTGGAAC AGGAAAGACC CTGATTGCTC TAAAGGAGAT GOTOGAACTG CCCCTGAGAC ATCCTGCCCT CTTTAAGGCA ATTGGTGTGA AGICCTIGAA IGAAGIAGOG TAIGAIGACA TIGGIGGCIG CAGGAAGCAG CIAGCICAGA

> 960 90 840 780 720 660 60 540

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GIGGGAIGCG IGCIGIGGAG TICAAAGIGG IGGAAACAGA ICCIAGCCCI TATIGCAIIG

AGCCGTACTT CCTGGAAGCG TATCGACCCA TCCGGAAAGG AGACATTTTT

crrorccord

TIGCICCAGA CACAGIGATC CACTOCGAAG GGGAGCCTAT CAAACGAGAG GATGAGGAAG

TOUTGOCCEAT TOATGACACA GIGGAAGGCA TIACIGGIAA TEICTICGAG GIATACCITH

TAGGGGATGT

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E SEQUENCE CHARACTERISTICS: (A) LENGTH: 3276 base pairs

(2) INFORMATION FOR SEQ ID NO: 291:

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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Œ. SEQUENCE DESCRIPTION: SEQ ID NO: 291

45 GCGACCGICG TITGAGICGI CGCIGCCGCI GCCGCIGCCA CIGCCACIGC CACCICGCG GCCGCTTGCG ATCAGGAGCC GOTTTOTICGE COCTGCTCGE CHACCGCCTG GAAGAGCCGA GECCCGGCCC ACCOTTOTTO COCCGACGCC TOGCTGCCGG TGGGAGGAAG CGAGAGGGAA

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55 50 8 CAGTOTTOCT GARAGGARAG ARGAGACGAG ARGCTOTTTO CATCOTCCTT TCTGATGATA TTCTCAAACA GAAGAACCGT CCCAATCGGT TAATTGTTGA TGAAGCCATC AATGAGGACA GAGAGGCGCG AGICOGICOC TIOCCACCOC ICGIAGCCGI TACCCGCGGG CCGCCACAGC CGCCGGCCGG ACAGTGTGGT GICCTIGICC CAGCCCAAGA TGGATGAAIT GCAGTIGITC CGAGGIGACA COCCATOGOT TOTOGAGOOG ATTOAAAAGG TGATGACOTA TOAACAGOOA

> 240 180 120

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480 420 360 300

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CCTCAAGGCT AACCTGCGCA AGTCCCCCAGT TGCCAAGGAT GTGGACTTGG AGTTCCTGGC

GATCAGCTCA TCTACATCCC ACTTCCTGAT GAGAAGTCCC GTGTTGCCAT

2220

2280

છ 35 5 8 CTCTACGOCG ATTTOGTCGC TITGACAGGG AGGTAGATAT TGGAATTCCT GATGCTACAG GAGCCAGAGT AACCCATCAG CACTGCGGGA AACCGTGGTA GAGGTGCCAC AGGTAACCTG CATTGATICC GAGGICATGA ACTICICIAGO AGITACIATO GATGACTICO GGIGGGCCIT GACGCTTAGA GATTCTTCAG ATCCATACCA AGAACATGAA GCTGGCAGAT GATGTGGACC GGAAGACATC GGGGGCCTAG AGGATGTCAA ACGTGAGCTA CAGGAGCTGG TCCAGTATCC AGAGGCTGCT CTGCAAGCCA TCCGCAAGAA GATGGATCTC ATTGACCTAG AGGATGAGAC TOGRACAGTA GCCAATGAGA CTCACOGGCA TOTOGGTOCT GACTTAGCAG CCCTOTOCTC TOTOGRACIAC CORGRERART TOCTGRAGTT TOGORTGROR COTTOCRAGG GRETTOTOTT

> 1500 1440 1380 1320 1260 1200 1140 1080 1020

1800

CTANGGACCT CCNGGCNGIG GGAAAACTTT GINGGCCAAA GCCAINGCTA ANGAANGCCA 1920 1860

1980

TOCTOACCGA GTCATCAACC AGATCCTGAC AGAAATOGAT GGCATGTCCA CAAAAAAAAAA TGATGAGCTG GATTCGATTG CCAAGGCTCG TGGAGGTAAC GOCCAATGIC AGAGAAAICI TIGACAAGGC CCGCCAAGGI GCCCCIGIG IGCIATICIT GOCCAACTIC ATCICCATCA AGGGICCTGA GCTGCTCACC ATGIGGTITG GOGAGICTGA ATTOGCOCTA CCAACCOGCC TGACATCATT GATCCTGCCA ATTOGRGATG GTOGTGGGGC 2160 2100 2040

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CTIGITCIGA TGAGAAGATT CGGATGAATA GAGTTGTTCG GAATAACCTT CGTGTACGCC

CATCAGCATC CAGCCATGCC CTGATGTGAA GTACGGCAAA CGTATCCATG

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2340 2400 2460 2520 2580 2640 2700 2760 2820 2880 2940 3000 3060 3120 3180 3240 3276

TAMANTGACT ANTGGCTTCT CTGGAGCTGA CCTGACAGAG ATTTGCCAGC GTGCTTGCAA GCTGGCCATC CGTGAATCCA TCGAGAGTGA GATTAGGCGA GAACGAGAGA GGCAGACAAA CCCATCAGCC ATGGAGGTAG AAGAGGATGA TCCAGTGCCT GAGATCCGTC GAGATCACTT IGAAGAAGCC ATGCGCTITIG CGCGCCGTTC TGTCAGTGAC AATGACATTC GGAAGTATGA GATGTITICCC CAGACCCTIC ACCAGAGICG GGCTITICGC AGCTICAGAT ICCCTICAGG GAACCAGGOT GGAGCTGGCC CCAGTCAGGG CAGTGGAGGC GGCACAGGTG GCAGTGTATA

S

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TOGGCTOCCT GOACCTTOTT CCCTOGGGGT GGGGGGCTT GCCCAGGAGA GGGACCAGG

CACAGAAGAC AATGATGATG ACCTGTATGG CTAAGTGGTG GTGGCCAGGG TGCAGTGAGC

15

STOCOCCCAC AGCTGCTCC ATTCTCCAGT CTGAACAGTT CAGCTACAGT CTGACTCTGG ACAGGGGGTT TCTGTTGCAA AATACAAAA CAAAAGCGAT AAAATAAAAG CGATTTTCAT

20

MOGIAGOCO GAGAGIGAAT TACCAACAGO GAATIGOGOCO TIGOOCIAIG CCATITICIGI TOTAGITITGS GECAGTICCAG GEGACCTIGTIG TOCOGTIGTIGA ACCAAGGCAC TACTGCCACC IGCCACAGIA AAGCATCTGC ACTIGACTCA ATGCTGCCCG AGCCCTCCCT TCCCCCTATC CAACCTGGGT AGGTGGGTAG GGCCCACAGT TGCTGGATGT TTATATAGAG AGTAGGTTGA TITATITITAC ATOCITITICA GITAATGITIG GAAAACTAAT CACAAGCAGT ITCIAAACCA raaaatgaca tottotaaaa ggacaataaa cottoooton aaatgoomra aaaaaaaaa

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S 120 180 240 300 360

PIGCANIGGE TGAAFTCCCC TCCTCACGCC AGCCTAGGAG AAGAAGTTCG TAGTCCCAGA GGTGAGGCAG GAGGCGGCAG TTTCTGGCGG GTGAGGGCGG AGCTGAAGTG ACAGCGGAGG

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SEQUENCE DESCRIPTION: SEQ ID NO: 292;

<u>z</u>

(A) LENGTH: 1695 base pairs

(1) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 292:

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STRANDEDNESS: double TYPE: nucleic acid

<u>e</u> <u>0</u> 0

TOPOLOGY: linear

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AAAAAAGGG GGCCCTCTA AAGNNCCANN CTTCGI

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CCTTGGCCGG AGGTTCGGGG ACCCCTTCGG CTGAAGCATT TGACTCGGTC TTGGGTCATA

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TCCGAGAAAA CAGCAACAAG CTGAGCTGCT GTGACAGAGG GGAACAAGAT GGCGGCGCG AAGGGAGCCT CTGGGTGAGG ACCCAACTGG GGCTCCCGCC GCTGCTGCTG CTGACCATGG

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COGAAGCAAC GSTCGGTGGG GCGGAGAAGG GGGCTGGCCC CAGGAGGAGG AGGAAACCTT

မ 20 5 35 25 5 8 55 S 25 8 CTOGTOGTOG GGGCGGCCG CATCGGCTGC GAGCTCCTCA AGAATCTCGT GCTCACCGGT TGANTGTGGT TCCGTTGGAC TGGGCTGAAG TACAAAGTCA AGGAGAAGAA ACGAATGCAT CAGATCAAGA GAAGCCGAAG CCAGAGCTAG AGCATCTAAT GAAGATOGTO ACATTAAACG TATTTCTACT GATGCTGATC AAGAAGTATC TCCTGACAGA GCTGACCCTG AAGCTGCCTG GGAACCAACG GAACCTATAC ATTIGCATOOT TIGGGCAAAG TACTIGITICA ACCAGITIGIT TIGGGGAAGAA ACAGCTOGGT ATCTTOGACA AGTAACTACT ATCAAAAAGG GTGTGACCGA GTGTTATGAG GCCCGAAACC ATGITAATAG AATGIGCCIG GCAGCIGAIG TICCICITAT IGAAAGIGGA TATAANGIGG AATTITINCCG ACAGITIATA CIGGITAIGA AIGCITIAGA TAACAGAGCI CTGCAGTTTT ACCCGAAAGC TAATATCGTT GCCTACCATG ACAGCATCAT GAACCCTGAC TITITIGITIC AAAAGAAACA TGITOGAAGA TCAAAGGCAC AGGITGCCAA GGAAAGIGTA TICICCCACA TCGACCIGAT TGATCIOGAT ACTATICATG TAAGCAACCI CAACAGACAG ATGGCACTOT COCGGGGCT GCCCCGGGAG CTGGCTGAGG CGGTGGCCGG GGGCCGGGTR GAATGAACCC CAGTTAGOCC TGAAAGACCA GCAGGTTCTA GATGTAAAGA GCTATGCACG AAGGAANGOG CTAAANCAAC NGGANANGAN CCAGINNAAA CTITITIACCA AGCITITIAA TOTCATCCTA AGCCGACCCA GAGAACCTTT CCTGGCTGTA CAATTCGTAA CACACCTTCA GGAAGGATTG AAGATTTTAT CAGGAAAAAT AGACCAGTGC AGAACAATTT TTTTGAATAA AGGGAACATT ATTECTGCTA TIGCTACTAC TAATGCAGTA ATTGCTGGGT TGATAGTATI CCTCAGGATG CATATTTTCA GTATGAATAT GAAGAGTAGA TTTGATATCA AATCAATGG TGAGCTCATA TOGGATAAGG ATGACCCATC TGCAATOGAT TITGTCACCT CTGCTGCAAA TCTTTTTTCA AAGAGCAICG AGACTTTGAG AGTTCATTTA GCAGAAAAAG GGGATGGAGG AGATGACATC AGGTATCTGT TGACAATGGA CAAACTATGG CGGAAAAGGA AACCTCCAKT TGACTTCCTC CAGGACTATA CTTTATTGAT CAACATCCTT CATAGTGAAG TAATAATCAC AAGAAGTTGT CAGAATTTGG AATTAGAAAT GGCAGCCGGC CCAAATTGAA GATGGGAAAG GAACAATCCT AATATCTTCC GAAGAGGGGAG ACAACCAAAC CCAAGAAAGA AGCTICTIGT GCCTIGTGCA CTGGATCCTC CCAACCCCAA TOTTCTCACC TTACAAGACA AGATAGTGAA AGAAAAATTT GCTATOGTAG CACCAGATGT TIGITAIGTA IGIGCCAGCA AGCCAGAGGI GACIGIOCGG CIGAAIGICC ATAAAGIGAC CCCGGGTGCN GATTIGGCAGN GCCTCCGCCG CGGCTCGTGG TTGTCCCGCC ACCTAGGAAA TTCAAGCAG AGACGGAAGC 1020 1680 1440 1380 1320 1260 1200 1140 1080 180 1620 1560 1500 360 300 240 120 960 900 840 780 720 660 9 540 480 420 8

1260 1200 1140 1080 1020

1500 1440 1380 1320

1501

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ

ID NO:

E SEQUENCE CHARACTERISTICS: (A) LENGTH: 2683 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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2 INFORMATION FOR SEQ ID NO: 294:

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25 8 . 45 6 3 30 . 5 5 S 80 TAGTAGATTT TATAAGCCAC AGAGACAAAC CAGAAACGGA ATAATGTTAC TITGGATGCT TTATGAACAA ATAGGATOCC TAGTIGAGGA TGITCCCAAA GTITTGTCCA ATCITATCAT TOTOTOAGIG ATAGAGTATG GGAGGGACOT COCTAGOTIG GAAAATGAGA ATIGAAGGG TOCHOTOGOC CTOCHAMATE AGACAGAAAT GGCTTGAGAA GCCGCAGGG AGCATGCCTG CCAACGCTCT TCTCTTATCC ATCTGTTTTG TIGITTATIC AAAAICCAAG ACACIAIGCC AAIGCAACCG IGACIACITI GOGAGAIIGG GCATCACTTA TCAGCTATOG TCAACCTGGT TTCATCTGTA TCTCTCTCTT TTCACCTGTA TIGCCITIAA AGICTIACGA CITICCCCAT TITAGICIAA IGGGAAGATA CAGAIGIGCA TRATTITIT CATTAGCATT GITATCAGCT TGTACTGGTC TCATAACTCT GGTTTTGGAA GAATAATTTG TTGAGACCCT TCACCAGAAT GTCAATCITT TITTCTGTGT AACATGGAAA CTTGTGTGAC TAGTETETT TGATGGTGAT AGTGATGGGG TGCACTATCA TAATCACATE AGGTETGETT TCAGACATOC ACAGAAGTOG AGAGGATGOT CCTTGGACCC MATGTGTCCA TCACCTAGCT AGICIOCITI TIIGITITITI GITAITAITT TITITITITI GCICIGIGIT AIGGACAITT ICITITICITA TACTIGGTAA TITGACACCC TGITAATAAC GCAATTATIT CIGIGITCIT AAACAGIATA GAAATTOTTO CTOTOTTCTO TGAAAATAAC CTCCCCAAAA TAATTAGTAA CTGGTTGTTC AACWAGAGAG AGACTGTTCT GTTGTAAAAC TCTTTCAAAA ATTCTGATAT GGTAAGGTAC ATTANCANGG AGTOTOMANA AGANATGAGA GGGATGOTTO CTTTROCCTT GCATOTACAA AATAGTTGTA AGTTTGCATG CATGATGGAA AAATAAAAAC CTGTATCTCT GTTAAAAAA TTICCTTITA AUGITAACTA AUGAAGITCC AGAGAUGGGC CITAGAAAIG TOITTIAAGA GCTAGGTGGG GGATTTGGAG CAAAACCGTC GAGTAGGCAT GATACTGGTA GITICIAGGIG IGGCITIGIA CAIGCAGAAG AAIGCIAIAI GCIGCACAIT GOGRAGOCCIC ATTOTANGIT COTCARGAGA GICCITIGGOT TARAGOTGIA CIGAGGGIGG GIAAGAAGGA CIGIAICIAC ACCIGITCIT CCCIACCIIC GWICTCCOGG TTATTTCCAG TOGGTGTAAA AGCAGAGCTG GGCCTTTCCC

> 960 900 840 780 720 660 600 540 480 420 360 300 240 180 120

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TAGTETECAA TATOGAGEAT CTEAAGETTE TECTOGGGGA TOGGGATTOG GATOGGEAGA

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240 480 540 909 999 720 780 840 900 960 1020 1140 1200 1260 1320 1380 1454 8 120 180 300 1080 1440 TOGOTOCTICC AGGOGCCCC TOCGCCCCAA GACACTCCTG, CTCACCAGCT CCGAGATCTT TGGACAGCAA TGGGCTGCAA TTAGGCAGAT AAAGACATCA GTCCCAGTAA ATGAATCCAT AGACTICATICT AGCACCAACT ACCATTAGCA CTATGTTAGG AGCTGCAAGG CCCCAAAGTA CONCORGAT GAGGACTIONS TECACTACCS ACTIGOCOGAG TITICCAAAG ÁGCOGCOGA GAGAGACAGG TACCGGCTGG ACGATGGCCG CCGCGTCCGG GACCTGGACC GAGTGCTCAT ATTCCTGCTG CCTGTATTCT CTATTCCAAT AAAGCAGAGT TTGACACCGW MAAAAAAA OGGETACEAG ACCTACEGE AGCETEACE CTEGITYTTEG ATGACGTGEA AGGTEATGAE CTCATTGGGCA GTOTCACCCT GGACCACTTT GGGGAGGTGC CAGGTGGCCC GGCTAGAGCC AGCCAGGGC GIGAAGICCA GIGGCAGGIG ITITGICCCCA GIGCIGAGAG CAGAGAGAAG TCACCGGCTA GCCCAGGCCA CAGCCAGCCT GTCGTGTCCA GCCTGACGCC TACTGGGGCA GOSCAGCAGG CITTIGIGIT CICTAAAAT GITTITAICCT CCCITIGGTA CCITAAITIG AGRANGAATG CCGGGCCCCT CAGGGCTGTT CGGTGTGCTG TCAGCCTCCC ACAGGTGGTA CAGCOSTIGCA CACCAGTISTIC STISTISTICACIO STISTITAACA CGTGACACTIG recrirente CHETHOCHER TECHETHECT STREETETTS SCATCINGCT GCTAATCCTG AGGCTGGTAG CAGAATIGCAC ATTIGGAAGCT CCCACCCCAT ATTIGTTCTTC AAAGTIGGAGG TCTCCCCTGA TECHGACAAG TOGGAGACC COTOGGGGA GGGACCTOG AGCTCCCAGC ACCAAGGGTG ACCETGGCAT GCCCCACAAA CAGATCACCA GCCAGCTTAC ACAGGCATTA ACTCTCCTCA atgaggaaga atcattcaca actgagcaag acattcatat gatcatttaa ggaagtgttt CCCTINIGIG TINGCANGIN TANICGGCIN ACTOCINAAT CCCNAIGAAT AGICCINGGC CICAICICGC IGITGGCICG CCAGIGGGAG GCCCIGIGIG GCCTGAGCTG CCTGTCGAGC ACTIGICCTICG CAGADATICTIC DACATIGIFIC TIGITICITIC TOTALITITIC TOTALITITICG TOGGICTICAC TITYTCTTCT ACACGTCCTT TCCTGAAGTG TCGAGTCCAG SEQUENCE DESCRIPTION: SEQ ID NO: 296: (A) LENGTH: 828 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 296: AAAAAAAAA AACN X; 3 8 S 9 15 20 22 8 35 <del>수</del> 45 S 55 1740 1800 1860 1920 1980 2040 2100 2160 2220 2280 2340 2400 2460 2520 2580 2640 9 120 180 240 300 360 420 2683 GGACCTICAA TITICAAGTIG TICGICATGC CCCCGAAAAA GIGGGGSCCA AACAAGCTGA AGATOCTOCC AAAAGCATAA CCAATOGGCA GTGATGATOG AGCTCAGCCC TCCACCTCCA CAGCTCAAGA GCAAGATGAC GTTCTCATAG TTGATTCGGA TGAAGAAGAT TCTTCAAATA ATGCCGACCT CATGAAGAAG AGAGAAGCCG CAAGAGGAAA TTAGATGAGA AAGAGAATCT CAGTGCAAAG AGGTCACGTA TAGAACAGAA GGAAGAGCTT GATGATGTCA TAGCATTAGA TTGAACAGAA ATGCCTCTAA ACAGAACCCT CTTACTATTT AGTTTATCTG GGCAGAACCA GATIGITATS TCCTITISTIC CAAAGGGAAA AAATTGACAG CAGTGACTTG AAAATGATTC TGCTCCCTTT GAAAGCATTC ATTITTGCTAG AACTGTTAGA CACATTGCAG TATGCTGTAT TGAAAGTAGG AATATAGTTT TAAAAACCCT TTGAACAAAG TGTGTGCATA ACCAGTCATG AGATAAAACA ACACAATGCA TOTTGCCTTT TTAATGTAAA TACCCTTAGG TATCATTAAT AGTITICAAAA TAITIGIGGIT TAGTAAAGIT GATACCIGGI TAIAAATAIT AIGCCITITAI GATGGCACTA TOTATATTAA TOTAAAACAA TOTTAATTTTA CTCAAGITTT CAGITTIOTAC ATATTCCATG TITTCTAAAA CICAAATAAC GACTATACTT ATGGACCAAA TAAATGGCAY GGACTCGGGG TGGCTCTAAG GGGCAGGGAT AGGGCTGGGG AGGGCGGGCC TGTGGCCCTG CTCTGGAAGG CACGCCCTICG CCGGAGCCTG TTGACAAGGA CTTCTACTCC GAGTTTGGGA ACAAGACCAC AGGGAAGATG GAGAACTACG AGCTGATCCA CTCTAGTCGC GTCAAGTTTA CCTACCCAG IGAGGAGGAG ATTGGGGACC TGACGTTCAC TOTGGCCCAA AAGATGGCTG AGCCAGAGAA GOCCCCAGCC CYCAGCATCC TOCTGTACGT GCAGGCCTTC CAGGTGGGCA TGCCACCCC TITITGCTAG AAGAAGAATT ATTITITAGCC TAGATCTAAC CATTITICATA CICTITAACIG ATTGAAACAG ATTCAAAGAA GTATCGAGTG CTATGCATTG AAACTTGTTT TTAAATGTTA COCCIGGIAT GICTOTOTAA GAAGCCAAIT ITTGTGTATT GTTACAGITT CAGGTTAITT CAGCTACCTG ACGCACTGCT TCCTCCAGCA CCTCATGGTC GTGCTGTCCT ACCAGCCCCT TCTCGTGCAG GTTCCACCCC GATGCAGGTG GTCAGGTGCT SEQUENCE DESCRIPTION: SEQ ID NO: 295: TGCATTCTKG TKAAAAAAN NACAGAAAAA AAAAAAAAA AGA (A) LENGTH: 1454 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDWESS: double
(D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 295:

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6 35 30 25 20 5 3 55 50 GOCAGAGTOT CCCAGCCAGA TOTOTGCCCC CACCCCATOT CCATTTACAT GICCTTCAAT GIGIATICAG AGCACCTYTC CAGAIGCACC ATGCATGCIC ACAGICCCIT GCCTAIGIGI ATCAGGGATC TTTTCACACT GCTGTTTTTT GCCCACCTCA ANAGGYACYT CTTCTGTANA GCTTTCCCTK GGTATCAGGA ATCANAATTA GAGAACAAGG CCAACCAAGG GACAGACTGG AAAGCACTTA GATGTTTAAG GAGGAGAAAG GATGGAAGCA ACCCCAAGGA TATGCAAGAA GGGCATGATG AACCCCCTTC CCTCTGGCAG OCTABACIGO TOTINGCITA GANTINIGOI TINCINGAGA TOTAGCAGAT AMGIOGGITA AAANCTAGAC AANAGAAGIC ANCGANAGCA GCITTITCCIC AAANGIGIGA CICCICAGGG TTTTTTCTTT GAGAAAGAAG TOGACTGGGG CACAACTTTT AGTCTGAGGG GAGCTAGTGG AACAAAACCT ACTATCACCA TCATCCTTCA ACAGCCACAG TCTGAATTGA GCCAACATTT TCAATTTCCA TTAACTCAGA TCAGCCATTG TGATTCACCA TTTGTCAGGC TCTCAGGTTT 3 TCTCATTCAG CCTTACAGCA TAACTAATTA TITGTITTCC TCACTACA CCTCTAGCGC AGTAACATTT GCAGAATTGC AGATTTTCCC CCAGATACTA GGAGGAAAGG ATCACTACCA TOOTGAACT AGTTATATAG CTTOCAGACA TGAGGGAGAC ATCAAACAGG GGGAAGCTTT GACCAGTCCT CATTOTOTOT TOCATAAAAC ACTATATTTT TITOGAAATG TTACTGTCCA AAAGCCTCTT TITITATITI GICATOCICI IGAAAANGIT IGACCATIIG TAGIATACAC AGIGAAACTI TGTTTTAAGA TAGGAAAAAA AAATAGTGGG CAAGGTGAAC ATCAGACGTA AATTTGTGTG INFORMATION FOR SEQ ID NO: 297: E Œ SEQUENCE CHARACTERISTICS: GGTGGGGAAG SEQUENCE DESCRIPTION: SEQ ID NO: 297: (A) LENGTH: 2416 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear GGGTCGTGGT GTTTTAAAAAG CATAAGTTAC CTGTTTGCAC TGCCTTTTGC CAAGITCAGC CAGITCTCCG CTGCTTGCAA CCICITIGGT CCITCTAICA CIAAAACICA 660 600 540 8 720 900 840 780

GAAGATGTGC ATAATGTCTG CTCTTGTGTA GCTCAGGAGA CAATTCCAGC ACAGACACTA ACCCCCAGG CCTGAAGCTG GCCCTTGAAG GATGGATGAA ATTTGGATAG AGAATGAGGA CAGTTAACGC TGAACTGCAG CTGCAAGTAA TAGCAMGAAC AGTCAGAAAA ATACCTTATG AGACAGAGGG NCTCCAAGTG AGAGAAGCAT GAAAAATGAG CARGGGCCTG GATCAGTGGG 828 780 720 660 600 540 480 420

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CAGTGAGGAG CCCAGGGATG GAGGGGGTTC CTGAAGTATT GCAGTTGGCT GTAGTAGCTG AGTICTITIC CAUGITACCG AAACIGTAGC CAGTTACAGT TTACTCAGGA AAACGGTAGA

AAATGAGOOT AACAGACCAG ACTGCAGCAA GTTATCAGAT TCCTCAATCA GATGCACTAG

AAGCCAGCOT GTGCCCTCTG GTTTAGTGAG TGTAATAGAG TCCCTGGCAC

TACCCATCCA AGACCTAGAG CATGAAACAG GGCCCTTTCC AAGTAGGCTC TGGGTGTCCT

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OCCAMAZITE AMBCETTOCE TITTIMBOMGG CEMECAGEMG ACCOMETTOG TECTECTTOT TCAATTCAGC CAIGGTAGTG CTGGTTGGCA GGGATTGGTA ACGGAGAGAA CTGCTCATCA

180 120

300 240

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CATCCCAGGG CTITOTGACA GICTCTAATI CCCTICCCTI CICGITAAGA AICATATIGI CTTCCTAGGA CACCCGAGCT GCTTGCCCAG GGTCCTGTTT CCCTGCTAAC TCCAGAGAAG GATGCTTOTG TAATCGACCA CCTAGCCTTC TCTCTCCCCT CCCGTCCTCC CCCAGAATCA CCAGCCCCTC CCATTGCCAT GGCAMAACAG GTACCTTTGG GGCATGGGGG CATTACATGG

2160 2100 2040

GAATTOTOAA GOTCOGAGIA OTTAGATCIT TAGCTITTAT TCCTTATITI TITOTATTAC ATAGTAGCTT TCAGACCATA CAGTATICAT TGGGTTACTC CTATTATTAT CAAGTAGCTG

TCTCCATGTG TATAAAITAT TGATCATGTT GCTGGCTTTT ATAAACTCTA AGCGAAGGAG

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GAGTETETGA GETTTTCAGT CCAAATETTT GCAAGGETCA AAATGCCACA GAACETETCC GGCTGCAAAR CGAGAAGAGA GCCGGTGGAG TGTACTTGTC CCTGACAGGC TGACCTACCT TOTOTTATAT TCTOCTTOTO AATAGCTOGA GCAAACCTOG GOCTGACACG CGTAAGSTAG TTATATATTO AGIGATGAAT TGATCCTCIT TITICCCTAA GGGATATGAA TIGITITITCT GIATTITICI GIICACAGIA TIIGIGIGIG IGCIIGITII GGCAGCICAI ITIGGCIGIA TGATAAGTGC TTTAAGCAAT GTCCATACCC CGTCAAGACT CCCAGCTTAG TCATTTTCTT GAGCAGIGAT ICICITITCI CICCCCACCC CCIGCCCITI GITACCAACA CCAGITICCC CTOGIACTTO TOTACATOCC GOTGACCTGA GGACTCCACT CACACTOOCG AGCAAAAAAGG

1740

1680 1620 1560 1500 1440 1380 1320 1260 1200 1140 1080 1020

TETTETECCE ACTECCEATG GEAGGGACCG GACCATECET ACATGGAACA TGCTGTTCCT

1980

1920 1860 1800

360 420

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GAGCACTICCC TCACCCTTTC

CACATGGTAA TGAAGCACTG TTTTTAAATA AAAGRGRGAA

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COCTOCOTTY COTTYTOOTA TOTACTICCY TOATACTIGG TITACTICATIC ACCORDIGADA

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INFORMATION FOR SEQ ID NO: 298:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 545 base pairs

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720 780 840

	(B) TYPE: nucleic acid (C) STRANDENNESS: double (D) TOPOLOGY: linear	
5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 298:	
	GAATTCGGCA CGAGCCATGC YTGGCCTCTC CTTGATTCTT ACAGTCACTT TGTTGGCTGT	09
9	TICTGACTCA GCAGCTACCT GCATTGTGGC CAAAGGATGA CCTATTCCTT CTCAGGAGGG	120
2	CANANATOTIC GANTAGTORY TOTCCATGCC TCTCCTCATG GGCTACCACC TCTGCCACG	180
	TOSTTANTCA GTACAACCA GAGAGAAGC TOCTGGAACT GACCTCTGGG AACTCCCTGG	240
15	ATGGTTTGGT GCAGGAATGT AGTAGGCATA CACGTGGTTG CGTGGATCTG GGCCCTCCTG	300
	ATCTGAGTAG AGAGGTAAAA GSSCACCATC TCCTTGACCT YTGGGGAACT CATCCACAAA	360
Ę	GAAGATOTTT CCAAGATGCT TCTGAAGATT GSCTAAAAAT AGCCGGTTTC CACCCCCGTG	420
3	AATGCATCCA TTCTAGAATG CTCCTTCACC AGGACCAGAG AACTGATTTA CAGAAGTGAC	480
	ATGAAAACAT TOCATOCCAG AATTIGGANT ACCTOAAATT NAATTITCTAC CTATTAAAA	540
25	NAAAA	545
30	(2) INFORMATION FOR SEQ ID NO: 299:	
35	(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1510 base pairs (B) TYPE: nucleic acid (C) STRANDELNESS: double (D) TOPOLOGY: linear	
	(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 299:	
9	GGCTCTGCTG GGCATCATAC TTGTCACTGG GTAAACAGTT TGCCCACTTA CCGCACATGA	09
	AGCTGCTTGC CAGGGCTCTC CGGCTCTGTG AGTTTGGGAG GCAGGCATCT TCCAGGAGGC	120
¥	ווסדופכרוסה בכאמסטאוסד סונסססטטטב מסכמאסססום כונסמכרוכככ סוככאמסוסם	180
}	TTGGGCCCAG GGCTGATCTC CCACCCTGTG GAGCCTGCAT TACTGGAAGG ATCATGCGGC	240
	CHGATTGATGC CAACGTGGCC GGCAATGTCC ACGGGGGGAC CATCCTGAAG ATGATCGAGG	300
20	AGGLAGOGGC CATCATCAGC ACCOGGCATT GCAACAGCCA GAAGGGGGAG CGCTGTGTGG	360
	CCCCCTCCC TCGTGTCGAG CGCACCGACT TCCTGTCTCC CATGTGCATC GGTGAGGTGG	420
2	COCATGTCAG COCOGAGATC ACCTACACCT CCAAGCACTC TOTOGAGOTO CAGGTCAACG	480
3	TGATGTCCGA AAACATCCTC ACAGGTCCCA AAAAGCTGAC CAATAAGGCC ACCCTGTGGT	540
	AIGTOCCCCT GTCGCTGAAG AATGTGGACA AGGTCCTCGA GGTGCCTCCT GTTGTGTAIT	009
9	CCCGGCANGA GCAGGAGGAG GAGGGCCGGA AGCGGTATGA AGCCCAGAAG CTCCAGGGCA	099

1020 1140 1380 900 960 1200 1260 1320 1080 1440 1500 1530 120 TOCACACCAA GTOCAGGAAC GOSCACATCO TCCAGCCAGT CCTCAACCCA GAGCCGAACA CIGCACGCCA CIGCAAGACC AACAICGICA CAGCTICCGT GGAGGCCAIT AAITITICATG CHETCAGCTA CAGCCAGTCC AGCTTGATCC ACCTGGTGGG GCCTTCAGAC TGCACCCTGC ACCCCTITIOT CCACCCACOT OTCACCATCA ACCTCATGCA TGAGGTCGCC GGGATCGTGG ACAAGATCAG AAAAGGCTGC GTCATCACCA TCTCGGGACG CATGACCTTC ACGAGCAATA AGICCATICAA GAICGAGGIG ITGGIGGACG CCGACCCTIGT TGTGGACAGC TCTCAGAAGC CITINGAAGITI CCCCCCITIGG CCAAAAACCC AATIICACATI GAGAGCTGGI GITIGICTGAA GITITICGIAT CACAGIGITA ACCIGIACIC ICTCCIGCAA ACCIACACAC CAAAGCITITA TITATATCAT TCCAGTATCA ATGCTACACA GTGTTGTCCC GAGGGCCGGG AGGCGTTGGG CAGAAACCCT CGGGAATGCT TCCGAGCACG CTGTAGGGTA TGGGAAGAAC CCAGCACCAC AGGINGTGAG AGACACATTA CACCTAACCA ACAAGAAGAA GGATCCTCCC CCTTATAATT GCTACCGGGC CGCCAGTGCC TTCTTTCACCT ACGTGTCGCT GAGCCAGGAA GGCAGGTCGC TOCCTOTOCC CCACCTOOTS CCCAACACC AGGACGAGAA GAAGCGCTTT GAGGAAGGCA AAGGGCGGTA CCTGCAGATG AAGGCGAAGC GACAGGGCCA CGCGGAGCCT CAGCCCTAGA CTCCCTCCTC CTGCCACTGG TGCCTCGAGT AGCCATGGCA ACGGGCCCAG TGTCCAGTCA TAACTATICIT TACAGGGAAT CCGTACATTC TGGCTTCCCG AGNATTTCGT CCAACATGTT (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 300: (1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 997 base pairs
(B) TYPES nucleic acid
(C) STRANDENESS: double
(D) TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 300: TMATAAAGCT GNTGCTTTGGC TGGGGAAGNA S 2 2 13 25 ಜ 32 45 6

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180 240 8 360 420 480

TIGANGAACC CINAAICCCA ACAACIGAIT GAAIGGGIAA AAGACACTIA IAGCCCAGAI GAACACCTCT GGGCCACCCT TCAGCGTGCA CGGTGGATGC CTGGCTCTGT TCCCAACCAC CCCAAGTACG ACATCACTTC TATTGCCAGG CTGGTCAAGT GGCAGGGTCA

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CAACAAGTIT GACCCAAAGG TAGATGATAA TOCTCTTCAG TGCTTAGAAG AATACCTACG

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TATCTGCGTT TATGGGGCTG GGGACTTGAA TTGGATGCTT CAAAACCATC ACCTGTTGGC

TEAGGGAGAC ATCGATAAGG GTGCTCCTTA TGCTCCCTGC TCTGGAATCC ACCAGCGGGC

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Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 2369 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

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(2) INFORMATION FOR SEQ ID NO: 302:

55 8 GGAGCTATTG AAAAGGCTGT AATTTGGCCT CAGTATGTCA AGGATAGAAT TCATTCCACC GOGGETOCTG TIGGICTIGG AGEATIGICE TACTATOSET ISSGACIGIC TAATGAGATT GAACCATCGA TOGAAAAAAT ATTTAAAATT GATCAGATOG GAAGATOGTT TOTTOCTOGA AMAICAMGAA TIOGGAITCCG GCGIOGGAGA ACIYGCCAAG AACICAAAGA GGCAGCAFIG GTGAAGAATT CCATCACGAA GAATCAATGG CTGTTAACAC CTAGCAGGGA ATATGCCACC TOTCTCCGGA CACTACCTIC TAGGGITTIC CACCCAGCIT TCACCAAGGC CICCCCTGIT CCAAAGCATC, TIGCTIGGIT GCTACATICI GGIGIGATGG GIGCAGIGGI GGCICCICIO ACCCCTOTIC TCATGAACTI CATGATGAGA GGCTCTTGGG TGACAATTGG TGTGACCTTT TATATOTACT TAGCAGGGAG TATTGGTTTA ACAGCTTTGT CTGCCATAGC AATCAGCAGA CATTICAGAT CIGCICGGIA GACCIGGIGC ACCACCACCA TGTIGGCIGC AAGGCIGGIG TTGAGGCCGA CGCTAGGGGC CCGGAAGRAA ACTGCGAGGC GAAGGTGACC GGGGACCGAG CIGGGAGING GCCIGGÓRCI CGICTITIGIG ICCICATIGG GAICIAIGIT ICTICCACCI GOCCICICCA CIGIGOCCAI GIGIGOCCC AGIGAAAAGI TICIGAACAI GOGIGCACCC GEAGCCATGG TIGGAGCIGG AATGCIGGIA CGAICAATAC CATATGACCA GAGCCCAGGC GGGGTCCTCT TCTCATCAGA GCTGCATGGT ACACAGCTGG CATTGTGGGA 360 300 240 180 120 540 480 420 660 8 900 780 720 60

30 25 ξ. SEQUENCE DESCRIPTION: SEQ ID NO: 301: (A) LENGTH: 2345 base paix
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

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Ξ SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2345 base pairs

ຣ INFORMATION FOR SEQ ID NO: 301:

CAAAGTAAGG AAAAARAAAA AAAGAAAAAA AACTCGA 960 997

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TIGATOGAAA GAGAACCIIC CCIICIGIAC IOTIAACITA AAAAIAAATA GCICCIGATI TYCCATYCTG TGGAGCTGCC GTTCCTAATA ATTCCAGGTT TGGTAGCGTG GAGGAGAACT GGGATATOTO GTAGAGCACT TGATTTCAGT TGAATGCCTG CTGGTAGCTT 900 840

CCTCCACTAA CTTTCTCACT AAGTGAGAAT GAGAACTGCT GTGATAGGGA 780

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GACTCAGCTT CIGOCTICIC TOCTACATICA AATATCTTOT TTAATOGGGC AGATATGCAT

TARATRISTIT GTACARICAG CTITICGITIGA AGTITIRGARG ATRAGRARICA TGICATICATR

CCGGTAATGT GATGCCTCAG GTCTGCCTTT TTTTCTGGAG AATAAATGCA

THIOGROTTC ARABATHIGA TECCHITRAC TEGRITOCIGA GIRICIACAT GGATREATTR AGCHTOTICC TICTOTATGA TACCCAGAAA GTARICAAGC GTOCAGAAGI AICACCAATO ACCACCOTOS CTOSTOCCAC TCTTTACTCA GTOSCAATOT ACGSTOGATT AGJTCTTTTC

AATATATTTA TOCCAGITOC AACTATOCTG GCAACTGGAG GCAACAGAAA GAAATGAAGT

1140 1080 1020

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TITAAAIGIT

GTAATCCTCT CCCAAATAAG CACACACATT TICAATICIC AIGITIGAGI GATITTAAAA

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CCTAGTAGTT COTTIGUAAAT TOCTIGOCTIGG GTGRATIGOTIG CITTGTTCTCT CACCCCTAAC 720

TICGIGGCAT CCITTAGGAT AAGAGGGCIG MIAITAGAIT GIGGGTAAGT 660

ATGTACAAAC ATGCTCAGAA CTTGCTGGGA CAGTGTGGGT GGGAGACCAG

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GGGCAAGAGC TTATAAGGCC ATCTATGGGA CTGAACTTTG AGACACACTA TGAGAGCGTT GCTACCTGTG 540 600

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TIMAMATITA GIAGOTICAC IGAGIAACTA AAATITIAGCA AACCIGIGIT IGCATATITI TOTTTTOOTG AATOTGAAAA CTAAAGTTTG TOTCATGAGA ATOTAAGICT TTTTTCTACT

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TITIOGROTICO AGRATIATIOT ARTIARITOTO ATRAGIGRITI TOGROCTITIG GIRARGOGRO

1560 1500 1440 1380 1320

CAGAGAGAAG GAGTCACCTG CAGTCTTTTG TTTTTTTAAA TACTTAGAAC TTAGCACTTG TOTTATTGAT TAGTGAGGAG CCAGTAAGAA ACATCTOOGT ATTTOGAAAC AAGTOGTCAT 1620

23

TOTTACATIC ATCTOCTOAA CITAACAAAA CIGIICATCC IGAAACAGGC ACAGGIGAIG 1680

CATICICCIO CIGINOCTIC ICAGIOCICI CITICCAATA TAGANGIGGI CANGITIGAC 1740 1800

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TTOTACAGAA TOTTAATCAT ACAGAGAATC CTTGATGGAA TTATATATOT GIOTTTTACT TTTGAATGIT ACAAAAGGAA ATAACTITAA AACTATICIC AAGAGAAAAT ATTCAAAGCA 1920 1860

TGAAATATOT TOCTTTTTICC AGAATACAAA CAGTATACTC ATGATTGCTA AGTOTTTTTT

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NATOCITICAT TAGITITOCCC TAGCAGACITI TIACITICICI

TACACTOCTA CACCATTACT

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AAGG G

TTCTTGAGAC

ATTIGIAAGI CCITIGATAC AGAAGAGITA TATTIAGGAG GNCTITAAIG

.2345 2340 2280 TATTITIOCA TATTIATICA ACIGICIAAT IGAATACAGC TIGCICTIGI CACCICIICA

2040 1980

TITITICICCI AAGACTITIG GITTCICGCA TIGCCICICA GACTAAGCAC TAAAAAGCAA AGCTITCAAG CCTTTATAGA AAAGCTICTT TOTGGCTTAC ACTGGAAAATT ATGAAAGCAG

2100 2160

ACCAAAACAG AACTAGTNCT GTCTTAATGA AATATATCAA CCCAAAAAGTG TAATGAGGAA 2220

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(D) TOPOLOGY: Linear		
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 302:		CAAGAACGGC TICCIGIACT ITCTIGATGA GGCGCTGCTT AATGTIGITA TIGGIGAGG
TITITITITI TITITITITI TITITINCAG ATCATIGITI ATTIATIACI TCAGATAAA	\$ 09	GATGTGTTGT CATGTCAAAA AGTAGGAAGT TCTGTTTCTC TGTTGTCAAT ACACCCTTTT
AGATAGTATA CATATTAGGG AATCCCTTAA AATTCAACTC TAGAGTTATA CACCATCTAG	120	CCACCAGGIT TITAGCIAAT CGITCCCGIA CATTICITAA CIGATAATGC AATTITAATG
TACTITIGCA ATGAATGITA ACAACAACA AAAAAATCIC TAAACACCTG AAGCCCCAC	081	GATTCCATGT CTCACCACTA AGTAATTCAA TCCAGTTCTG GACGGTTTCT GGAGGCTGAA
TATTAACATG GACTATGGTA ATAAAAATT TTGACATTTA ATFTGTTCAA CATATAGTAT	240	TITICCTIAAC ATGCTTCAGA GCTTCATCAA GAAGAACATC CCCTGTTGGA GCATCTGACT
THACATTATG ARACCHATGG TGATGATACA ATANGTGAT ARAGNATAG TANANITAIA	300	TACAGATTAC CTITICITISTE AATAGACITT TACGICICAE TCCACAAGCC TCTAGTISTA
CITIAAAAAG CAAGGITTA TAGTCTGACA ATGCTAATTA TCCTAATTGT ATATAAAAA	360	ACCTTCCTCT CANTECTANT TCANTTAACA TACAGCCACG TAATCCAGAT GATATACAGT
TIAAAACATA GAGCITICTG TIACAAAATT CTIAATCCTC 1936T15TAA TCATTACTTG	420	CATTCCAAAA TGATGTGTAA ACCTTCGCGG TCCTTGAGGC CCAGCAGGAG CACTTCCTCC
CTACCAATIT ACATGCAACA TCTGCTAGGA CTGACATITG ATTITITICC CCAAGAATGT	480	ATCAGGGTCA GCCGCGTTTC CTTGGAGTCG CCCTTGTCGT CGTCGTCCTG CTCGTCGCG
STGAGTAGAT AAATGACATT TCAGAGCAGA TAITAATTTA CITISTICGACA GAAAAAGAAA	540	CASCITCTECS CATCATCTC GCTGCTAGCC GCGCCGCCCC CCGCCGCCCC CTCCTTGTCG
CTCAAGHTTG GTACTGGTCA CAAGCCTCTT CCCAATAGAA AITHATAAAA CAGTAAGATA	009	acaecantae acanacene acrocacea
AAATTTAAAA AAAATCTAAA AAGGGGATGC ATAGGCAAAG AGTACCATAA ATGGCACAGC	660 25	
TCAAAAATC CCAGGACCAA TCAGACACAC ATCTTTTCTC TCTCCTTCAG GGACAAGAGG	720	(2) INFORMATION FOR SEO ID NO: 303:
TOGATTITICC CATCAAATAA CCATGATTGA ACCAAGOGAG GGGCACCAGG TSTACAACTG	780	
ATTAGATICTT GCAAAATACT AAGATGGGAG CAGGGGGC CAGAAGAAGG GGTAATTTAT	840	į
ATATAATTCA AACTATATAC AGCATAAATG GAATGCAGCC CATCCCAAAC TGGCTCTGTG	006	
AAACAATTGG ACCTTTATAG TTAAAATTAT AACAAGTGTA ATAATACAAT AGATTTACAT	35	(xi) SEQUEN
GOGAAGCAAA ATCCAAGGGA CATTTTATAT TAAGTATTA CTGTGCTGTT TCAATTTAAA	1020	GODACOTOTO GITTICACTIC GIGOSCCTICC CCGTGGGTTT GCGACGTTTA GCGACTATTC
AATAATTITIG CIAAGIAIAC AICTCAACIG AAGICTAIGI AAAAAAIGIC CIAAIAGAIA	1080	_
CAGATATTTA CCTTTGGTGA GTTGAAGGCC TTTTTGGGAC TTCTGTCTGA ACTGTAGGCA	1140	GOGGICCCT GGGCTCCAGG CIGTIGCGGG GIGTAGGTGG GAGTCAGGA CGGTTCGGGG
GARTOCTAGA TOTACATGCA CATATGGAGA AACTCAAGCT GAGGTCATCC AAAAGCTGTG	1200	CCCCHACTOT CCCCBARGT GCCCACATG GCCGCAGGG GAGAGCATGG CTCAGCGGAT
COTATGAGGA GEOTGGAGGT ACTITGAAAG TCAAAGTAGA CCAGAAACCC AAACAGGTA	1260	GGTCTGGGTG GACCTGGAGA TGACAGGATT GGACATTGAG AAGGACCAGA TTATTGAGAT
ACAGTGAGGA TGGCAACAGG GAATGGAATG CCAATATGGC AGTAAAACTT TTTTTAAAA	1320	GOCCIGICIS ATAACIGACT CTGATCTCAA CATTITIGGCT GAAGSTOCTA ACCIGATTAT
CAGANAGAGO ANGOCTICTIC GTACCAGCAG ANTOCTISTAC ACGTACAAAA AAGAAAAAGC	1380 50	AAAACAACCA GATGAGTTGC TGGACAGCAT GTCAGATTGG TGTAAGGAGC ATCAGGGGAA
CACCCACCAT TITGIAAAAC AGAAGCCAAT TATAGTGTGG GAAAGTACAA ATTACAGAAA	1440	GICTGGCCTT ACCAAGGCAG TGAAGGAGAG TACAATTACA TTGCAGCAGG CAGAGTATGA
ACCAGAROTC AACAGAAGAA AAACTACTGG TTTTACTTGAG AGAAAGGAGA ATGGTTCACC	1500	ATTICTIONCE TITIONAGAE AGRAACTEC TECAGGGCTE TOTECACTTG CAGGAAATTE
CCGAGCAGAG TTACTTIGGTG AACOCCGCCA CCACCCCCA CAGAACCTCA TTGGTGTTCG	1560 55	AGITCAIGAA GATAAGAAGI TICTIGACAA ATACATGCCC CAGITCATGA AACATCTICA
CCTICAGACA TICCACITICA GGOTCIANGI CGAGARRITIG CCGCACTICIC TIGGIAGCCA	1620	TIATAGATA ATTGATGTGA GCACTGTTAA AGAACTGTGC AGAGGCTGGT ATCCAGAAGA
ANTENTACTO CTOSTOCAGA AGAGGAGAA AAGCATTOTO CAGGACGTOC GAGGCATGAG	1680 60	ATATGAATTT GCACCAAAGA AGGTGGTTC TCATAGGGCA CTTGATGACA TTAGTGAAAG

1800 1860 1920 2340 2369 1740 1980 2100 2220 240 540 909 2040 2160 2280 120 180 300 360 420 480 CCAGGTAAAT GAGGGCCAGC AAGCGCCTGT CCATGCGGTG AGGGTCATTC ACCCATTTGT CACCAGGIT TITAGCIAAT CGITCCCGIA CATTICITAA CIGAIAAIGC AAITITAAIG PAGAACIGC TICCIGIACT TICTICATGA GGCGCTGCTT AATGITGITA TIGGIGAGG INTICCATOT CTCACCACTA AGTANITICAA TCCAGITICTG GACCGITITCT GGAGGCTGAG TICCITIAAC ATGCITICAGA GCITICATCAA GAAGAACATC CCCTGITIGGA GCAITCIGACT ACAGATTAC CITICITOTI AATAGACTITI TAGGICICAT ICCACAAGCC ICTAGITGIA CCTTCCTCT CAATGCTAAT TCAATTAACA TACAGCCACG TAATCCAGAT GATATACAGT NITICCAAAA TGATGTGTAA ACCITCGCGG TCCTTGAGGC CCAGCAGGAG CACTTCCTCC ICAGGATCA GCCGCGTITIC CITIGAAGICG CCCITIGICGT CGTCGTCCTG CTCGTCGCG BACTICTIBGS COTOSTICCTC GCTGCTAGCC GCGCCGCCC CCGCCGCCCC CTCCTTGTCG GACGICIG GITTCACCIC GIGOSCCICC CCGIGOGITT GCGACGITTA GCGACTAITG CCGAGGTGT CCGCGAAGGT GGCGCACATG GGCGGCAGGG GAGAGCATGG CTCAGCGGAT OCCIGICIO AIAACIGACI CIGAICICAA CAITITIGGCI GAAGGICCIA ACCIGAITAI AAACAACCA GATGAGTTGC TGGACAGCAT GTCAGATTGG TGTAAGGAGC ATCACGGGAA ITCTGGCCTT ACCAAGGCAG TGAAGGAGAG TACAATTACA TTGCAGCAGG CAGAGTATGA ITTICTIONEC TITIONACIAC AGCAGACTCC TCCAGGGCTC TOTCCACTTG CAGGAAATTC GITCATGAA GATAAGAAGT TTCTTGACAA ATACATGCCC CAGTTCATGA AACATCTTCA AIGIGITICI CAIGICAAAA AGIAGGAAGI ICIGITIICIC IGIIGICAAI ACACCCIITTI OCCIOCOCC ACOCCOGCTO COAGACTGGG GCCGTGGYTG CTGGTCCCGG GTGATGCTAG COGCICCCT GOGCICCAGG CIGITGCGGG GIGIAGGIGG GAGICACGGA CGGIICGGGG CICTICICITIC GACCTICEAGA TCACAGGATT GGACATTICAG AAGGACCAGA TTATTICAGAT (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 303: (1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1181 base pairs
(B) TYPE: nucleic acid
(C) STRANDERNESS: double
(D) TOPOLOGY: linear (2) INFORMATION FOR SEQ ID NO: 303: COGCOTTOC GOGAGOCCTC GOTOCOCCO

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GAGGAAAATT ATAGAAAATG GGGAAAATGA GAAGACCGTG AGTTGATGCC AGTTATCATG

ARCGITARCE GOAGGCAACE TOIGGIGGIT TITTITITCIC ACGCIGATGG

900 840 780

CATCHARGAG CTTCAGTTTT ACCGARATRA CATCTTCHAG ARABARTRG ATGRARAGAR

20 30 25 2 Ö 3 6 35 50 8 55 GIGNTAGAGG GGATICCTOGA CACACTOGAG GGCCCCAACA TCCCGCCCAT CCAGAGGGTC COCAMOSTAG CASTOTROST GICTICOGACC GIGOGCCCCA CACAGCGGCT GCTCCTCTGT GCAGCCAATG CAGAAAATGG GTCAGCTCCT TTGAGAAACCC CTCCCCACCT ACCCCTTCCT ACCTOTOGGE ACTICIGAMA GEACAAGGEE AMGAACTECT GECLAGGACT GEAMGGETET TOTTICOTAG ATGGGGTTTG CAGCIGCCAC TGAGCIOTAG CIGCGIAAGI ACCICCITGN AGTECTETTT CECGEAGETG CAGAGAGGAG GAAGACTATT AAAGGACAGT CETGATGACA ACAGCCAAAG CTGTTGCGGT GACCCTGCAG TCACACTGAC CCCACCTGAA AFTCTTGGCC COCAGAGALA TOCOTROCAT GOTOCOTROCT COTROGOTTO COACCACOT COTCAACGOO GOCACCCIGO CIGCCCIACA CAIGCICITIC CIGCICIAIC IGCAITITIGC CIACCACAAA TITAATICCC TTACCIACIC TIGCICIAIT TITTIATIIG AAATGGAGAT GAGCAAAATA 2 AAAAAAAAA AAAAAAAAAAA AAAAAAGGG GGNCCCC TOCTOTTIAT CICTOCCACA TIGICTIGCT AAATATAGAC TIGGTAATTA AAAAAAAAA TTATATTTCT ATTCACATCA TTATTGAAAA TACCCAGCTC AGTGCCTGGC TTAATAAATG GGATGATCCT GAGCIGITCA AACAAGCIGT ATATAAACAG ACAATGAAAC TCTTTGCAGA TOCATOCCAA AACCAATOCC TOCCAAACAA AATCTTAGAC ATCCCAATAT AATATOTTAG GAGGGGCTTG GTTTAAAAAA AGACCTTTCC CTCTCCCTGC CCCTAGAACA ACCAGTATTA GAAGAAAGAG AAGAAAAATC GGACCTTCCT GAGACCACCG AAAGTAAAAA TGGAGCAACG TOAGGAAAGT COAGATGOTC GTGTGGACAG CTGGCGAAAAC TTCCAAGCCA ATACGAAGGG GGAAGAAGAG ATTGAAGCTC AAGAAAAAGC CAAACGGGAA AGAGAGTGGC AGAAAAAACTT GCTGGAAATT AAAAGGAAAAG AGAGAGAAGC CAAAGAGATG CATGAAAGGA AACGACAAAG ACACATTCAT GGCTGAAGCA ATTITITIGGA CATTICITGT TACCAAAAGA TCTATAATCA CGAAGGACTC ATTCTTTCCT TEAGTBACCG CCCAAGGTCA CAGGCACAGA ACCTTTCCCC TOCATITION TITICAATACG INFORMATION FOR SEQ ID NO: 305: E SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 1493 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear AFTTAATATC GATCAGAGTA AFTCTTTTGT CCCACTTCCA CCCCAACATA GAGTAGTATT TGCTTTTTAG TOCTATORIC CITECTICITY ACATTGAAAI 1500 1440 1380 1320 1260 1200 1140 1080 1020 1537 960 360 300 240 180 120 420 600 540 480 780 720 660 8

20 5 25 5 6 330 ٠ 45 8 55 50 CGAPATRICTA TITTICTCCT ARTRIGCTOT TICCATTRIG ACACAGCAGC TCCTTIGTAA GTACCAGGIC ATGICCATCC CTTGGTACAT ATATGCATIT GCTTTTAAAC CATTICTTTT алдаалааа аалаалаала алааалаааа аалаааааа N CTTGGCAGAG CAGGCTTGGGA AAAGACAAGC CTCCAGGGCC TTCAGCTTGT ACGCCAACAT CGACATCCTC GAGGAAGAAG TAGACGCTGA TGCAGCTGAT GCAGCTGCTG CTGAAGAGGA GGATGGAGAG GOGACATOGO AACTACAGOG GOGOCGGOGG GOGGOCGCCG AAAAATOGAGO TGGCCCGGAA CCACATETTO CCCACTCCGC GCGCGGGGCT AGCGCGGGTT TCAGCGACGG GAGCCCTCAA CITITITISTOT TECCOCECANT CECACETETE CIECAACECTG GACGICTACE TICEGGAGGE 2 TICCIOGGCA TGAAGGGCIT TAAGGGACAG CIGAGCCGGC AGGIGGCAGA ICAGAIGIGG ATGGATGATA CCTCAGGCTC TAGCTTCGAG GATATGGGTG AGCTGCATCA GCGCCTGCGC CTGGTCTTCA CTCTGGTTGC TATCCTACTC CATGGGATGA AGACGTCTGA CACTATTATC CCTATICAAGA TOGTICAACTIT CCCCCAGAAA ATTIGCAGGTIG AACTICTATIGG ACCTICTICATIG AGACCCTACT TIGATOTOGA GCCTGCTCAG GTGCGAACAG GGCTCCTGGA GTCCATGATC TOGOGROGOT TOGRAGRARA CATOCROGGO GGROGOTORG CTOTGRITTGR CATOGRGARA ATCCACCICC ACOCCOPOTT CTACCICTIC TOGOTOTIOG TOGGIGGACT GICCACACIG TCATCCTTCA COGGAGGCA CCCTGATGGG CACAGCCATT INFORMATION FOR SEQ ID NO: 304: (i) SEQUENCE CHARACTERISTICS: Ĕ CHCTTCGGTT AACTTGCATC TCCAGATTGA TTACTCAAGC AGACAGCACA TITACTICCT TGCCTACCTG TGCAACGCCC AGATCACCAT SEQUENCE DESCRIPTION: SEQ ID NO: 304: TOGGCTATIGG CCTCTTTGGG CATTGCATTG TCCTGTTCAT CACCTATAAT (A) LENGTH: 1537 base pairs(B) TYPE: nucleic acid(C) STRANDEDNESS: double TOPOLOGY: linear GGCACCIGCT TCGGCTACTG GCTGGGAGTC GCTGCAGATG 1140 1181 1080 1020 960 540 480 420 360 300 180 120 780 720 660 600 900

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700 0711	Š
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TGACCGAGAG ACCAGAACCT GCTTGCTGGA GCTTAGTGCT CAGAGCTGGG COGCOGCICC TCTGCTGTCA GCGCCGGCAG CCCCTCCCGG CTTCACTTCC CETISETACTIC AGAAGETICCG GGATCCCAGC AGCCGCCACG CCCTGGCCTTC GCTTCCAGTC AGGCCAACAC CGACGCGCAC TGGGCAGGAA GACAGGACCC CATCTGCACA GAGGTCCTGG CTGGAACCGA GCAGCCTCCT CCTCCTAGGA

TGACAGGGGA GAGTTAAGCT CCCGTTCTCC ACCGTGCCGG CTGGCCAGGT

240 300 360 420 480 540

CTCCAGCTCT CCAGTTTTCA GGTTGGAGAC ATTAGATGGA GGCCAAGAAG GECEGACAGA GGAAAGCTGG ATTTTGGGAG CGGGCTGCCT CCCATGGAGT GOSCOAGGAC COGAAATTCG CCCCTTCAGA TAAGAGTCAA CCTCCAACTA

9 9 720 780 840 900

TCCCGGGGTG TCCCCGAGGA TCTGGCTGGA CTTCCAGAGT ACCTGAGGAA TACCTCACCG ACTTCGGAAA TACACAGAGG GCTCCACAGG TAAGACGGCC TOTOCTGAAA CCTTAAGGAC GGGGTCAATG CCTGCATTCT GCCACTGCTG GOGACTICTIGG CAATCCTCAG CCCCTGGTAA ATGCCCAGTG CACAGATGAC GCCACAGGGG TCTGCACATC GCCATTGAGA AAGAGGAGTC TGCAGTGTGT GTGGAGAATG GGGCCAATGT GCATGCCCGG GTCTGCGGCG ACTTCTTCCA

ACAGGIGCCA GICAGCGGA TCCAAACCGA ITTGACGGAG AICGGCTCTT

120

CTCTCNCAAT ATGGCTCCCC CGGGCTGGCA GRWRKTCRGT CWCKRGTGGC

	GAAGSTOCCA CCATTGGTGC TGCCTTCTCT TCCCACAGCC TGTAACTCAG TGTTTTTGTAC	840	(2) INFORMATION FOR SEQ ID NO: 307:
v	TTCACTGAAT TOTCATGOTT AGAVACTTCG TOGATAGTTT OTGGAAATCA TCCAATTAAA	900	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 2860 base pairs
<b>.</b>	CATACTECTT AAAACAGTOT TGCTOTGACT TCAGAGACAA GCCTOGAAGG GGCACCTTAG	096	(B) TYPE: nucleic acid (C) STRANDEDNESS: double
	GAAGCCCCTT CGCTTCAGTT GCTCGCTTCT GGCTGTGCTC CCTTCGAAGG CCCAGATAAG	1020	
01	ACAGGGAACA CTITOTGAGCA CACAGAGCAG CATCTGATGC CCTTOTGGTGT TTGGCATGTG	. 1080	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 307:
	CCCCCTOTCT ACTRACCART CAGTOTOCCA TGAGGCCAC GCCACCCAAA CCTTTCACTT	1140	GIGINGACCG CICICNCAAF ATGGCTCCCC CGGGCTGGCA GRWRK
<u>.</u>	TOCAMBRIC TRACCOTICCT CCACCORGIA CCATOTICCTA GCCTOTICTO ATTIGITACT	1200 : 15	TAGCCTOTCC TGACAGGGA GAGTTAAGCT CCCGTTCTCC ACCGT
2	GOTAMINITE TITINIGIATA ATAMITITI ATACCCAAGE CAITGAIGIA CITITICCTIG	1260	GGCCTGAGGG TGACCGAGAG ACCAGAACCT GCTTGCTGGA GCTTA
	TACTICICCET TISTISOSTICCE TISTICINGCT TISCINGAACC CCAAANTICET TISSOSTINGS	1320	GAGGGAGTT COGCOGOTOC TOTACTOTOA GCGCCGGCAG CCCOT
20	ACAGACCTIGG CTGAACCTTA GTTTCTTCAT CTATGAAATG GGAATATGAA TTACTGCAGC	1380 20	TOCCOCAGO COTOCTACTO AGAAGTICOG GGATOCCAGO AGOOG
	ACCTITIAGG GCAGATITISC CATGGCATAT ACAAGGTAAC TACCATAGTS CTCCTTGGCT	1440	AGCTIGGGG GCTTCCAGTC AGCCAACAC CGACCGCAC TGGGG
;	ATTGCCAATA TCCTATTATT TCTGTGTAAA ATGAAGATAC TGATTGTTTT GAG	1493	TTGACATOTO CATOTGCACA GAGGTOCTGG CTGGAACCGA GCAGO
3			TGACCTCACC CTCCAGCTCT CCAGTTTTCA GGTTGGAGAC ATTAG
			ATGGCTCTGA GGCGGACAGA GGAAAGCTCG ATTTTTGGGAG CGGGC
30	(2) INFORMATION FOR SEQ ID NO: 306:	30	CACAGTICCA GGGCGAGGAC CGGAAATICG CCCCTICAGA TAAGA
	(1) SPQUENCE CHARACTERISTICS: (A) LENGTH: 577 base pairs		CCBAAGGSA ACAGGISCCA GTCAGCCGGA TECAAACGGA TTTGA
	(B) TYPE: nucleic acid (C) STRANDEDNESS: double	:	CANTIGOGOTO TOCOGGGOTO TOCOGGGGA TOTGGCTGGA CITTOD
35	TOPOLOGY: line	35	GACCAGCAAG TACCTCACCG ACTTCGGAAA TACACAGAGG GCTCC
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 306:		TCHIGAAGGC TGTGCTGAAA CCTTAAGGAC GGGGTCAATG CCTGC
5	AATTCGGCAG AGGINITATA TACACTATAC TGGCATTTAC TGTTTCACCC AGCCCGGAAA	60 40	CAGATCGACC GGGACTCTGG CAATCCTCAG CCCCTGGTAA ATGCC
?	GICAGAGATO TATATIGGAA AATTTACAAC TCCANCTACA TIGGITCCCA GGAGGCICIC	120	TATTACCGAG GCCACAGGG TCTGCACATC GCCATTGAGA AAGAG
	ATAGCACATT ACCCAAGAAT CTACAAGGAT GATAAGAACA CCTATATTOG TTATGAACTT	180	GAAGCTCCTG GTGGAGAATG GGGCCAATGT GCATGCCCGG GTCTG
45	GACTATATCT TATAATTTTA TIGTTTATTT TGTGTTTAAT GCACAGCTAC TTCACACCTT	240 45	GAAGGSCCAA GGGACTIGCT TITATTICGG IGAGCIACCC CTCTC
	AAACTTOCTT TGATTTGGTG ATGTAAACTT TTAAACATTG CAGATCAGTG TAGAACTGGT	300	CAAGCAGTGG GATGTGGTAA GCTACCTCCT GGAGAACCCA CACCA
20	CATAGAGGAA GAGCTAGAAA TOCAGTAGCA TGATTTTTAA ATAACCTOTC TITGTTTTTG	360 50	GGCCACTGAC TCCCAGGGCA ACACAGTCCT GCATGCCCTA GTGGA
	ATCTTANACA GTANTICCCA GTAGTGACCA AGANCACAGT GATTATATAC ACTATACTGG	420	CAGCIGAGAA CATTGCACTG GTGACCAGCA TGTATGATGG GCTCC
	AGGGAPTICA TITITAATIC AICTITAGA AGATTIAGAA CICAFICCTI GIGTITAAAG	480	SCCYTCTOCC CTACCOTOCA GCTTGAGGAC ATCCOCAACC TGCAG
22	ggaatottta attgagaaat aaacatttot gwacaaatg ytaaaaaaa aaaaaaaa	540 55	AAGCTGGCCG CCAAGATCGAG ATTITCAGGC ACATC
	adadadada adadadada adadadada adctoga	577	TCAGGACTGA GCCACCTTTC CCGAAAGTTC ACCGAGTGGT GCTAR

1380 1440 1500

TEGETISTATIC ACCTOSISTIC TOTIGACAGE TSTEAGGAGA ACTEAGTICST GEAGATEATT

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960 1020 1080 1140 1200 1260 1320

> GOGACTIGCT TITATITICGG TGAGCTACCC CTCTCTTTGG CCGCTTGCAC GAIGIGGIAA GCIACCICCI GGAGAACCCA CACCAGCCG CCAGCCIGCA TECCAGGGCA ACACAGTECT GCATGCCTA GTGGATGATC TEGGACAACT CATTGCACTG GTGACCAGCA TGTATGATGG GCTCCTCCAA GCTKGGGSCC CTACCOTIGCA GCTTGAGGAC ATCCGCAACC TGCAGGATCT CACGCCTCTG CCAAGGAGGG CAAGATCGAG ATTTTCAGGC ACATCCTGCA GCGGGAGTTT GCCACCTTTC CCGAAAGTTC ACCGAGTGGT GCTATGGGCC TGTCCGGGTG

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OCCUTICATI GENAGAGECE GENECGACAE EGANIGOTES TITIOGRAGE CETGAACAA

1620 1560 80

CTGATCTACA TOTTCATCTT CACCGCTGTT GCCTACCATC AGCCTACCCT GAAGAAGCAG CTOCTOCAGG CGAAATGGGA TCTGCTCATC CCCAAGTTCT TCTTAAACTT CCTGTGTAAT TECHGETECT CCAGTECAAC TGATGGCCCA GATGCAGCAG GAGGCCAGAG GACAGAGCAG ATCTOGRAGO TOCHGRAGAGC CATCTCTGTC CTGGRGATGG AGRATGGCTA TOGAGETETT CANATTENCE ATCOGENTOG GEGNGETOGE ETTECHOGNG CHOCTOCHET AGGAGGCTTG GCGCCCCGAA GCTCCTACAG GCCCCAATGC CACAGAGTCA GTGCAGCCCA GTICCAGCAC ACAGGCAGIC TACAGITICA IGMICCCIGA AGCCCIGGIG AGCCIGAGCC COTOCOTORIO TOTOCOCTOG TOGOTOGGOOT GGCTGAACOT GCTTTACTAA TACACGTGGC COCTOCTION CAGTOGIGIC COAGGIGCIG TOTITICCIGG GCCATCGAGI GOTACCIGCO CONOTICATO TOGATOTOGT TOATAGACAG CTACTITOGA AATCOTOTIC CIGITOCAG ATCCTGCTAG GEOGEOCETE ACCIGAAAGE GGAGGITOGA AACICEATGE IGCIGACGGG CEACAICEIT TECTOSETTE CECTECEAAG GAGGATGAGG ATGGTGCCTC TGAGGAAAAC TATGTGCCCG TOCCTACOCT GTOTOLAGGAC CCGTCAGGGG CAGGTGTCCC TCGAACTCTC GAGAACCCTG CGATGAGCGC TOGTICCTTCA GGGTGGAGGA GGTGAACTGG GCTTCATGGG GAGCAGACGC TCAACATGCT CATCGCCCTC ATGAAGCGAA COTCACAGTG TCGCCACTGA CAGCTGGAGC TECGEGGEAT GETGETGETG CTGCTGCTGG CETALGTGCT GETCACCTAC ATCETGCTGC ATATTTICAC TAACTCAAAA AAAAAAAAAA AAAAAAAAA AAAANGAGG GGGCCCGKT AGGATETTIC CAACCACATE IGCIGGETET GGGGICCCAG IGAATICIGG IGGCAAATAT AGGAAAAAGC AGCGGGCAGG TGTGATGCTG ACCGTTGGCA CTAAGCCCAG TOCIACICIACIA GEAGGIACCIAG GGCIAACGGGG CCCAQTIACIAG GGGTIATCCTG GIAAGCCTCCT ASCCAAMTIC GCCCTATAAG IGAGIGCCWA TIACGATAAA (2) INFORMATION FOR SEQ ID NO: 308: 3 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 308: GOOGGATCTA CCTCCTCGTG GOGCCAGCTG TGGTACTTCT GGCGGCGCCA SEQUENCE CHARACTERISTICS: (A) LENGTH: 876 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (D) TOPOLOGY: linear ATGGCAGCCC Tracracrac 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2220 2160 2100 2040 1980 1920 1860 1800 1740 1680 2860

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SEQUENCE DESCRIPTION: SEQ ID NO:

60

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear (A) LENGTH: 2025 base pairs

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GETOCEGEE CEGGAETETO ATETETOTAG TOGECECETE CTECEGGGEE CETTITEGGE TOCTOCTOGO AACASOCCOO OCCCAAGTCC ATOGAGGTCT CCTOCTAGGC GGCCTGCCCA CACCCGCAGC COCTCGCACA CCAGCGAGGG GGCCCACCTG GACATCACCC CCAACTCGGG CADOCOCAGO COCTOCOADA COAGOGAGOS CACOCGAAGO COCTOCOACA COAGOGAGOG CATGACCCGC CTGATGCGAT CCCGCACAGC CTCTGGTTCC AGCGTCACTT CTCTGGATGG

ĕ 240 180 120

CCCTGCCTGC CATACTGCGC CTAACTCGGT ATTAATCCAA AGCTTATTTT GTAAGAGTGA

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COGTIGOCAC TOTOCTOCCO TOCCTCAGAG ACACCAAAACT GCCAAAAAACA AGACGCGTAC

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480 420 360

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GCTCTGGTGG AGACAAATGA GGTCTATTAC GTGGGTGCCC

TCTCCAAAGG CGGGGTGGCG

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SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 309:

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ઝ 25 20 ᅜ 5 TIBOCICCCC GAACCICCIA CAGOCICCAA TOCCACAGAG ICAGIGCAGC CCATIGGAGGG CIGCITOTOT CIGCOCIGGI GCIGGGCIGG CIGAACCIGC ITTACIATAC ACGIGGCITC GGAAGCTGCA GAAAGCCATC TCTGTCCTGG AGATGGAGAA TGGCTATTGG TGGTGCAGGA CATEGRACIE CIECUSCIEC TESCCIACOS ECIECICACO FACASCOSE TECTICAACAT CTICAAATIC ACCATCGGCA TOOGCGAGCI GGCCTICCAG GAGCAGCIGC ACTICCCCGG ACAGGAGGAC GAGGGCAACG GGGCCCAGTA CAGGGGTATC CTGGAAGCCT CCTTGGAGCT CAGCACACAG GCATCTACAG TOTCATGATC CAGAAGCCCT GOTGAGCCTG AGCCAGGANN CTICCCCICC CAAGGAGGAT GAGGATOGTG CCTCTGAGGA AAACTATGTG CCCGTCCAGC AGCGCTGGTG CTTCAGGGTG GAGGAGGTGA ACTGGGCTTC ATGGGAGCAG ACGCTGCCTA OCTOATOGOO CTOATGUAGO GAGACOGMOA ACAGTOTOGO CACTGACAGO TOGAGCATOT тсастаамим алааллаала алаалаалаа Астсба ITICCAACCA CAICIOCIGG CICIGGGGIC CCAGIGAATI CIGGIGGCAA ATATATATIT TCCTCCAGTC CAACTGATGG CCCAGATGCA GCAGGAGGCC AGAGGACAGA GCAGAGGATC AGAAGCAGCO OGCAGGTOTO ATGCTGACCO TTOGCACTAA GCCAGATOGC AGCCCCGATG GGACCCGTCA GGGGCAGGTG TCCCTCGAAC TCTCGAGAAC CCTGTCCTGG 840 660 600 540 480 420 360 300 240 876 780 720 180 120 8

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180 340 300 360 420 480 540

(2) INPORMATION FOR SEQ ID NO: 310: 55

(A) LENGTH: 3026 base pairs (B) TYPE: nucleic acid (C) STRANDEDNESS: double (i) SEQUENCE CHARACTERISTICS:

8

1620 1560

1680

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<u>, .</u>

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(D) TOPOLOGY: linear

TAGGCAGCAC TGAAATATCC TAACCCCCTA AGCTCCAGGT GCCCTGTGGN ACGAGCAACT GGACTATAGC AGGCTGGGC TCTGTCTTCC TGGTCATAGG CTCACTCTTT CCCCCAAATC TCATTCCCTT CCTACCCTCT CTAATGGCCC CTCCATTTAT TTGACTAAAG CATCACAAG IGGCACTAGC ATTATACCAA GAGTATGAGA AATACAGTGC TTTATGGCTC TAACATTACT TCATTACCTC CCTATCAGTT CTAGCATAGT AAACTACGGT ACCAGTGTTA GTGGGAAGAG AAGTOTAAAA AAAAGTCTTA ACAACAGCTT CTTGCTTGTA AAAATATGTA TTATACATCT ITCCICIGGA GCITIGGAGC CAAGGIGCIA AAAGGAATAG GIAGGAGACC ICITCIAICI AATCCTTAAA AGCATAATGT TGAACATTCA TTCAACAGCT GATGCCCTAT AACCCCTGCC IGGATITICIT CCIAITAGGC TATAAGAAGI AGGAAGATCT TTACATAAIT CAGAGIGGIT OCCITICAGIA TCAAGGCTGC CTGGAGAAG GATGGCAGCC TCAGGGCTTC CTTATGTCCT CCACCACAAA AGCTCCTTGA TGAAGGTCAT CTTTTTCCCC TATCCTGTTC TTCCCCTCCC COCTCCTAAT GETACGTGGG TACCCAGGCT GGTTCTTGGG CTAGGTAGTG GGGACCAAGT CHOCONTING CHANDANCO CACHGCATCO TACTOCTACO TOGICAACOO GCIGOTITOCA GGTATICGGAC CTICCTAAGTIG TGGAATTTACC TGATAAGGGA GAGGGAAATA CAAGGAGGC CICTOSTIST CCTOSCCTCA GCCAGCTGCC CACAAGCCAT AAACCAATAA AACAAGAATA CIGAGICAGI ITITIAICIG GGIICICITIC AITCCCACIG CACITIGGIGC IGCITITGGI GACTGGGAAC ACCCCATAAC TACAGAGTCT GACAGGAAGA CTGGAGACTG TCCACTTCTA GCTCGGAACT TACTGTGTAA ATAAACTTTC AGAACTGCTA CCATGAAGTG AAAATGCCAC ATTITICCITT ATAAITICTA CCCAIGITIGG GAAAAACIGG CITITITICCCA GCCCITITCCA GOSCATANAA CTCAACCCCT TCGATAGCAA GTCCCATCAG CCTATTATTT TTTTAAAGAA AACTIGCACT IGITITICIT TITACAGITA CITCCITICCT GCCCCAAAAT TATAAACTCT GINITITIAN AITICIGCICC IGANAAAIGA CIGICCCAIT CICCACICAC IGCAITIIGGG GCCTTTCCCA TTGGTCTGCA TGTCTTTAT CATTGCAGGC CAGTGGACAG AGGGAGAAGG GAGAACAGGG GTCGCCAACA CTTGTGTTGC TTTCTGACTG ATCCTGAACA AGAAAGAGTA ACACTGAGGC GCTCGCTCCC ATGCACAACT CTCCAAAACA CTTATCCTCC TGCAAGAGTG GOCTITICCAG GGICTITIACT GGGAAGCAGT TAAGCCCCCT CCTCACCCCT TCCTTITITIC ITICITITACI CCITIGGCIT CAAAGAITI IGGAAAAGAA ACAATAIGCI ITACACICAI TITCAAITIC IAAAITIIGCA GGGGAIACIG AAAAAIACGG CAGGIGGCCI AAGGCIGCIG (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 310: Ś 으 15 20 23 2 33 <del>6</del> 45 S 55

960 1020 1080 1140 1200 1260 1320 1380 1440 1500

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999 720 780 840

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45 35 25 55 50 CCTTAAACTO ACGCTACAMA ATAAACCTGG GCAAGTGAGG CAAGAGAAAT GAGGAAGAAT CTOGRIGACAG TRACATTITCA TIRACCAMAG AMBOTGGOTC ACCIGACCIC TGAAGAGCIG TITOGCCTOTC AGAGCTIGAT TAGAAGCCAA GACAGTOGCA GCAAAGGAAG ACTITOGCCCA CACCCAAGGT CANCCAAACA ACTIGGITGT GAACCCAACT GCCTTAACCT ICIGGGGAG AGATEATTET ACAAGTAATT TIGCACAGAE ATETECTEAC ECCAGTGEET GIETGGAGET ATTIATAAAT TIGAAAICCA AACTACTITC TIAATATCAC TIIGGICICC ATTITICCCA AAAAGATTCA AAGCTCTAAT AGAGTCACAG CTTTCCCAGG TATAAAACCT AAAATTAAGA AATTCTAAAA GAGAAGGGAG CTGAGGCCAT TCCCTGTAGG AGTAAAGATA AAAGGATAGG TAATGICCIT TCCCTGGAGT CAGIITITITT AAAAAGITAA CICTTAGIII TIACIIGIII AGTACCIGIA AGCATITIAG GICCCAGAAT GGAAAAAAA AICAGCIAIT GGIAAIATAA AAAGGAGATC ATTTAGITGG GTCTGAAAGG AAAAGTCTTT GCTATCCGAC ATGTACTGCT CCATCTOTGA GGTGAYAGGC AAGGATGAAA GACAAAGAAG GAAAAGAGTA TCAAAGGCAG AGTACTCAGG CCACTCCAAT CACCCTACAA GATGCCAAGG ARAGRACART AGRACTOGIC TICCRITITIG CCRCCITICC IGITCRIGRC ROCTRCIRAC TAAAGTTGAG GGGAGAGGAA ATCTTAAGAT TACAAGATAA AAAACGAATC CCCTAAACAA 2 AATTINCCAA ATAGAGATNG TATTAC AUGTAAAACA GAATATICIG TAAACCTAAT GICIGIATAA AIAAIGAGCG TIAACACAGI GOGTOCATOG AGGAATTOGG ACCTOGTTAT GTTOTTATTC TCGGACTGTG AATTTTOGTG GGAAAAACCT GTGGGTTOTG CTAATTTCTG TCCAGAAAAT AGGGTGGACA GAAGCTTGTG GGACAGGAAA TATGTCCCCC CCTAACTTTC TTGCTTCAAA AATTAAAATC CAGCATCCCA AGTACAATAA GCAGAGGTGG AAAATGATCT AGTTCCTGAT AGCTACCCAC AGAGCAAGTG AAAATATTCA ATAAGAAGTC AAAAAAAAAAA AAAAAAAACT CGAGGGGGGG CCCGGTACCC GGGGATTAGC TAGACTAGGA GACCCAGAAG TGAATGGGAA AGGGTGAGGA CTTCACAATG INFORMATION FOR SEQ ID NO: 311: (X) Ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 311: 909 € ) TYPE: nucleic acid
) STRANDEDNESS: double
) TOPOLOGY: linear LENGTH: 712 base pairs 543 AGGTCCCAGG AAGTCCAGCT PCT/US98/04493 2820 2760 2700 2640 2580 2520 2460 2400 2340 2280 2160 2100 2040 1980 1920 1860 1740 3026 3000 2940 2880 2220 1800

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GCAGOCTTTG TOCTCACCTA CAAGCTGGGT GAGCAGGGTG CCAGCAGCCT GTTTCCTCTT

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25 15 30 20 5 45 8 35 8 S 50 GOOTCAGCCT TOCTGAGCCT ATGTCTGCAG CACTTCTTGG GAGGCCTGGT CACCACAGTC CIBCIBCCIC IGITEARGIC GGIOCIBCGC TICCOCCICG GGGGCCIAGC CIGICAGACT GIBGICIBET CCATEBEIGG CICCICCTB GOIGGGACCT IGCIGGCCAA GCACIBGAAA CICCIBCIDG ACCACOOCOT TICIBCICCC GAGTIGGGAC IGIGGAAIGG IGIGGGIGCT CHOSCIGATE GOTTOGGGCC ACATECETEC TECTTECTEC TOCTCATECT CTCTGCCTTT GCCTTGGTCT TCCACCTGGA CACCCTGGGG GCCAGCATGG ACGCTGGCAC AATCTTGAGA GITATIGGGI TITIGGITGG TITITIGITIG TITITITACTA IGCITIGGIC IGIAAAAATA TIGCCAACCT ICTGGTTGAG CTGCAAGAAA ATATTTATGG TGAGAACTTT CAAAATTICA GAACTIICAG GAGGGCAAGA GAATATCAAA CAAAGATIIC TGGAAGTATT 3 AACTOGAGOG GOOGCCCOGT ACCCAAATCG CCGGATATGA TCGTAAACAA TC CONSTITUTOR ACCIDIDACOR AGCACCUAGO ACCITITUTOS GAGOIGAGIG GOIGGAGIGG ACAGCCTTCT ACCTICACIG GGAIGAIGCG CIGCAGCCAG CIGGCCCCCA GGGCCIGCAG GCCACACACT TTAGACTIAT TAATATGTIC TTGTCCTGTA TTTATACATA TGTGTATTIT GGAAAGTATT ACATRATICT GATGAGGAAA AAAAATCTIT GCAATTCTIT GCCTTTTTTA AGGGAAGCTA TAATICGATA CATAGIGAAA AAGGGAATGG TGACCCCTTT TGTGTTGTTT TCCCTGGATG AMATATCAG TATTAAGTAG ACAGCATATT ATTCAAGTGT TOCAACTGAA CTACATTCAG AAGGAAATAT TOTCTACATA GAATATTATA TGAAGTTGGT GAGGGCTCTC AATCAGCAGG GCCCCAGGAG GGAAGAAGAA GTGGGGCAAA GCCTGGCCTC TCTTATCTTG TTATTACGGT TTTATTAATT TTGTAGAGGG ACAGGGAGTG GGCAAGGGG GIGCCICTIC CACIGAGGAT AACAAACAGC ATIGTAATCC ATICTCITIGC ACCITCITCI AGAAGCAGCT TATTTGACTA ACCAGCCCCT CTGTGGTCCA CCAGCGTCTT GGCTTGGTGG INFORMATION FOR SEQ ID NO: 312: Ξ £ GGCCACGCTG GAGCTGCTGG GGAAGCTGCT GCTGGGCACT CTGCGGAGGC SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 312: (B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear ٤ STRANDEDNESS: double LENGTH: 1289 base pairs AMGCCATATT GITGITTITC Tererrice 712 540 480 420 360 300 240 660 600 180 120 300 240 180 120 720 660 600 540 480 420 360 6

	COX/III AAC	֚֭֭֚֚֚֚֡֝֝֝֝֜֝֜֝֜֜֝֜֜֜֝֜֜֜֜֜֜֜֜֜֜֜֜֝֜֜֜֜֝֓֜֜֜֜֝֡֓֜֝֝֓֜֜֜֝֡֡֡֩	
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	OCCOCTICAGO ACCITIVACIA TOTANGCIAC OCCTICETICA GOCCATACTE CITICAACITIS	780	65 70 75 80
	GANATOTCAA COGGNOCCCT TACACCAGOC CTCCAGCATC TAATAGACTT GAATCTACTC	840	Leu Gly Pro Glu Pro Lys His Leu Ala Leu Leu Pro Pro Arg Gly Glr 90
ν	TAAAGGAATA TITTAATCCAA CCTCACTACA TIGTAGCTCA GTCCAAGGAC TAACCCTGAA	900	or car man als car car Lau Pro Gly Gln Gly Pro Leu Pro Leu
	ATGROGOTOT TECHGEETTE AGGRANTG CEAAGOGOTE EECTGGGGGC TETGGCAGG	096	110
9	GOCTHATICS TOTOTOTING CAACCTITICG GTOCGACCTC CTOCGCCCCC ATGCGGTGAC	0707	Pro His Ile Asn Cys Thr Val Phe Ser Leu Lys Ala Ser Phe Ile Lys
3	CCCOTCOSTG TCTGTGTCTG TCCATAGGTG TGAGTCCAGC TAAAAAGACA AAACAGAACC	1080	į
	сотвоссска сстсевалься тесетовлея лектессаль втеслессет	1140	
15	GOGATGSCAT TOCGITGIGT GOCTTAITCC TGGAGAATCT GIATACGGCT CGCCTAIAGA	1200	
	aatatageet citcatgetg tattaaaagg actititaaa gcaaaaaaa aaaaaaaa	1260	(2) INFORMATION FOR SEQ ID NO: 315:
20	CTTGAGGGGG GGNCCGGTAC CCAATTATC	20	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 28 amino acids  (B) TYPE: amino acid  (D) TODOLOGY: linear  (*) SEQUENCY: linear
Š	(2) INPORMATION POR SEQ ID NO: 313:	\$6	und the letter yet let let let let let let let let let Ala Let
3	(i) SEQUENCE CHARACTERISTICS:	3	1 5 10 15
30	(A) LENGTH: 22 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 313:	30	Cys Asp Val Pro Ile Ser Leu Gly Phe Ser Pro Ser 20
	Met Phe Leu Ile Phe Val Tyr Phe Leu Lys Ile Leu Phe Ser Ser Ser 1 5 15		(2) INFORMATION FOR SEQ ID NO: 316:
35	Leu Pro Phe Leu Trp Leu 20	35	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 64 amino acids (B) TYPE: amino acid (D) TOPOLOSY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 316:
4	(2) INPORMATION FOR SEQ ID NO: 314:	40	Met Asp Gly Phe Ser Ser Arg Leu Phe Ser Ser Leu Pro Phe Val Al
	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 128 amino acids	,	1 5 10 13 10 12 12 10 12 12 12 12 12 12 12 12 12 12 12 12 12
45	(B) TYPE: amino acid (D) TOPOLOGY: linear	45	Leu Gin Trp Phe Lie Val Lie Sein his deu deu dei des des 20 20 20 20 20 20 20 20 20 20 20 20 20
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 314:		Ala Cys Cys Tyr Gln Thr His Cys Ser Leu Xaa Gln Leu Ser Ser Al
	Met Met Phe Leu Thx Gln Gly Gly Pro Leu Pro Ser Thx Arg		35 40 . 45
20	Gly Ala Leu Pro Lys Pro Ser Gly Leu L 25	90	Phe Ser Xaa Met Gly Glu Ser Cys Val Gly Glu Arg Glu Tyr Xaa Ph 50 50
55	Val Thr Cys Trp Asn Gly Leu Lys Gly Pro Leu Cys Gly Asn Arg Cys 35	55	
	Ser Pro Asn Thr Leu Leu Leu Leu Ala Ala Arg Gln Ala Leu Trp Lys Gly 50 60		(2) INFORMATION FOR SEQ ID NO: 317:
8	Arg Gly Arg Thr His Gln Asp Leu Pro Gly Pro Leu Gln Gly Arg Gln	09	(1) SEQUENCE CHARACTERISTICS:

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Lys Gln Asp Lys Lys 20 Met Pro Leu Ile Asn Leu Leu Leu Leu Tyr Tyr Val Pro Asn Gly Gly
1 10 15 Met Leu Trp Trp Ser Arg Asp Tyr Thr Met Val Phe Leu Leu Phe Thr 1 15 Pro Gln Gly Lys Lys Lys Lys 35 Met Gly Arg His Leu Val Leu Val Met Phe Ile Thr Thr Ser Leu His 1 10 15 (2) INFORMATION FOR SEQ ID NO: 318: Met Val Phe Thr Gly Asp Leu Val Ile Arg Gly Arg Thr Glu Leu Ser  $20 \ \ 25$ (2) INFORMATION FOR SEQ ID NO: 320: INFORMATION FOR SEQ ID NO: 319: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 318: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 317: (i) SEQUENCE CHARACTERISTICS: (i) SEQUENCE CHARACTERISTICS: (i) .SEQUENCE CHARACTERISTICS: (X Ĕ. SEQUENCE DESCRIPTION: SEQ ID NO: 319: (B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 320: (A) LENGTH: 21 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 33 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENOTH: 39 amino acids (A) LENGTH: 88 amino acids

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SEQUENCE DESCRIPTION: SEQ ID NO: 322:

(D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 27 amino acids

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Met Thr Phe Thr Leu Gly Asp Ser Gln Val Leu Leu Ile Asn Leu Phe
1 15

Pro Ser Met

Pro Ser Gly Ser Cys Ala Arg Pro 20 25

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(2) INFORMATION FOR SEQ ID NO: 323:

(1) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 64 amino acids

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(2) INFORMATION FOR SEQ ID NO: 322:

(i) SEQUENCE CHARACTERISTICS:

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Pro Gln Ser Pro Ser Leu Ser 20

Met Gln Pro Gly Ala Gly Val Leu Val Leu Gly Leu Leu Leu Pro Pro
1 10 15

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SEQUENCE DESCRIPTION: SEQ ID NO: 321:

(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 321:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 23 amino acids

5

Gln Thr Ser Glu Pro Ser Gly Thr

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Glu Thr Lys Gly Val Ala Leu Pro Glu Thr Met Lys Asp Ala Glu Asn 50 55

Leu Gly Arg Lys Ala Lys Pro Lys Glu Asn Thr Ile Tyr Leu Lys Val 65 70 75 80

Phe Ala Vai Leu Gly Leu Leu Ala Ala Gly Val Thr Leu Leu Leu Pro 35

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Met Val Cys Ser Ser Leu Cys Asp Ile Gly Gly Ile Ile Thr Pro Phe

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Ile Val Phe Arg Leu Arg Glu Val Trp Gln Ala Leu Pro Leu Ile Leu  $20 \ 20 \ 25$ 

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Met Cys Leu Glu Cys Trp Ala Glu Asn Leu Gly Pro His His Thr Ser

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SEQUENCE DESCRIPTION: SEQ ID NO: 323:

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Pro Ser Ile Pro Ala Met Phe Pro Asn Lys Ser Leu Leu His Cys Ile Phe Xaa 5 Ser Ser Gly Cys Phe Gln Gln Gln Glu Met 35 40 2 Leu His Phe Val 55 Ser Leu Leu Asn Pro Arg His Leu Cys Leu Phe Val Ser 50 ۲a Val 9

INFORMATION FOR SEQ ID NO: 324:

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SEQUENCE DESCRIPTION: SEQ ID NO: 324. (X ೫

Leu Ser Thr Ser Glu Tyr Ser Gln Ser Pro Lys Met Glu Ser Leu 5 Het. 22

Ser Ser His Arg Ile Asp Glu Asp Gly Glu Asn Thr Gln Ile Glu Asp 20 Pro Ala Glu Pro Met Ser Pro Val Leu Asn Ser Lys Phe Val 35 40 G). 돮

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Ser Ile Leu Met Asn Pro Ala Gln Asp Gly Glu Val Gln Leu 55 Asn Asp S 35

Lys Thr Lys Gly Asp Asp Thr Asp Thr Arg Asp 70 75 80 Asp Ile Ser Ile Leu Ala Thr Gly Cys Lys Gly Arg Glu Glu Thr Val 90 95 Gln Asn Asp Asp Ser 65 各

Leu Thr Cys Asp Ser Gly Ser Gln Ala 105 Ala Glu Glu Val Cys Ile Asp 100

Val Pro Ser Pro Ala Thr Arg Ser Glu Ala Leu Ser Ser Val Leu Asp 115 45

Gly Ser Glu Val Glu Glu Ile Pro Glu Thr Pro Cys Glu Ser Gln 150 150 Glu Ala Met Glu Ile Lys Glu His His Pro Glu Glu Gly 9er 135 130 130 20

Glu Glu Leu Lys Glu Glu Asn Met Glu Ser Val Pro Leu His Leu 175 Ser Leu Thr Glu Thr Gln Ser Gln Gly Leu Cys Leu Arg Arg His Pro 180 55

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Lys Lys 195 Lys Lys

(2) INFORMATION FOR SEQ ID NO: 325:

(D) TOPOLOGY: linear SEQUENCE DESCRIPTION: SEQ ID NO: 325: (A) LENGTH: 252 amino acids (i) SEQUENCE CHARACTERISTICS: TYPE: amino acid

Met Gly Gly Asp Leu Val Leu Gly Leu Gly Ala Leu Arg Arg Lys

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Arg Leu Leu Glu Glu Lys Ser Leu Ala Gly Trp Ala Leu Val Leu 20 Ala Xaa Xaa Gly Ile Gly Leu Met Val Leu His Ala Glu Met Leu Trp 45 2 2

Leu Ser Asp Phe Gly Gly Cys Ser Ala Val Asn Ala Thr Gly His 50

Leu Trp Leu lle Pro Ile Thr Phe Leu Thr Ile Gly Tyr Gly Asp Val 65 22

Va. Met Gly Val Cys Cys Thr Ala Leu Leu Val Ala Val Val Ala Arg Lys 100 Thr Gly 95 ž ile Val Cys Leu . 90 ξ Met Trp Gly Val Pro Gly Thr

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ABp Leu Glu Phe Asn Lys Ala Glu Lys His Val His Asn Phe Met Met 115 33

lle Gln Tyr Thr Lys Glu Met Lys Glu Ser Ala Ala Arg Val Leu Gln 130 Glu Ala Trp Met Phe Tyr Lys His Thr Arg Arg Lys Glu Ser His Ala 145

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Ala Arg Xaa His Gln Arg Xaa Leu Leu Ala Ala Ile Asn Ala Phe Arg 175

45

Gln Val Arg Leu Lys His Arg Lys Leu Arg Glu Gln Val Asn Ser Met 180

Val Asp Ile Ser Lys Met His Met' Ile Leu Tyr Asp Leu Gln Gln Asn 205 Ser Ser His Arg Ala Leu Glu Lys Gln Ile Asp Thr Leu Ala 215 Ser 210 Ę 8

Gly Lys Leu Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala Leu Gly Pro 225 235

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Ser Lys Xae 250

Arg Gln Leu Pro Glu Pro Ser Gln Gln 245

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45 55 50 Glu Lys Thr Thr Glu Asn Lys Glu Ser Asn Pro Phe Ile Leu Gln Val 50 55 Ile Val Ser Cys Arg Lys Asp Gly Pro Asp Ala Ala Ala Ala Pro Ala 20 25 Met Trp Arg Cys Arg Gly Lys Leu Ser Phe Pro Leu Phe Ala Val Val
1 10 15 Thr Val His Thr Cys His His Gln Ala Phe Leu Val Leu Ile Gly Trp \$35\$Ala Cys Trp Glu Gly Val His Ser Glu Pro Val Cys Arg Thr Val His 20Met Gly Glu Gly Lys Asn Gly Phe Gly Gly Phe Val His Thr Ala Asp 1 15 Asn Lys Leu Xaa 65 Val Ile Lys Asn Asn Ser His Tyr Gln Thr Ser Lys Ala Leu Glu Leu 35 40 Ser Lys Ser Gly Lys Glu Arg Lys Glu Ala Phe Leu Thr Ala Ile Ile 50 55 (2) INFORMATION FOR SEQ ID NO: 327: Lou Agn Ser Arg Ser Ile His Ile Ser Cye Ser Trp Pro Pro Ser Pro 65 70 75 80 Val Pro Gln Xaa 2 INFORMATION FOR SEQ ID NO: 328: 3 ξ (1) SEQUENÇE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 327: (A) LENGTH: 84 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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2 INFORMATION FOR SEQ ID NO: 326: (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 326: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 68 amino acids

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SEQUENCE DESCRIPTION: SEQ ID NO: 329:

(A) LENGTH: 63 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 329:

(i) SEQUENCE CHARACTERISTICS:

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Ile Asn Asn

Asn Ile Ile Leu Phe Leu Lys Lys Lys Ser Leu Phe Phe 20 25 30

Ile Asp Ser Val 35

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Met Leu Leu Ile Asn Leu Leu Trp Leu Val Thr Met Ile Lys Ser Val 1 15

(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 328:

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Asp Met Thr Val Ile Leu Arg Gly Arg Ala Gln His Lys Thr Ala Met \$35\$

Leu Ile Pro Gly Glu Ser Arg Leu Ala Pro Thr Phe Asn Pro Ser Ala
20 25 30

Met Thr Phe Pro Phe Glu Lys Lys Ile Val Ala Phe Ser Ala Phe Tyr 1 15

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Leu Glu Ser Tyr Asn Trp Lys Val Ser Cys Gln Leu Arg Glu Xaa 50 SS 60

(2) INFORMATION FOR SEQ ID NO: 330:

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Œ. (1) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 330: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 35 amino acids

Met His Ser Lys Gly Ser Ser Leu Leu Leu Phe Leu Pro Gln Leu Ile 1 15

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Leu Ile Leu Pro Val Cys Ala His Leu His Glu Glu Leu Asn Cys Cys
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Phe His Arg 35

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(2) INFORMATION FOR SEQ ID NO: 331:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 amino acids

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(A) LENGTH: 36 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 331: (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Gly Ala Leu Val Leu Leu Leu Cys Leu Leu Val Gly Val Gln Gln 1 15 15

Ser Gly Ser Val Trp Asp Ser 20

2

(2) INFORMATION FOR SEQ ID NO: 332:

(A) LENGTH: 40 amino acids SEQUENCE CHARACTERISTICS:

15

TYPE: amino acid

SEQUENCE DESCRIPTION: SEQ ID NO: 332: (D) TOPOLOGY: linear

Met Gln Ser Ala Glu Ile Leu Ser Trp Thr Asp Val Leu His Asp Phe 1 5 20

Phe Leu Trp Pro Ala Phe Glu Asp Arg Ala Leu Leu 25 3 2 Leu Phe Ser 25

Ile Phe Thr Leu Asn Gln Ile Val 35

23

(2) INFORMATION FOR SEQ ID NO: 333:

(A) LENGTH: 111 amino acids (B) TYPE: amino acid (i) SEQUENCE CHARACTERISTICS: 35

SEQUENCE DESCRIPTION: SEQ ID NO: 333: (D) TOPOLOGY: linear

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Met Gln Ser Leu Val Gln Trp Gly Leu Asp Ser Tyr Asp Tyr Leu Gln 1 5 15 6

Gly Leu Ile Gly Leu Leu Leu Ana Arg Gly Ser Lys Ile Lys Lys Leu 35 45

Asn Ala Pro Pro Gly Phe Phe Pro Arg Leu Gly Val Ile Gly Phe Ala 20

Val Tyr Pro Pro Gly Phe Met Gly Leu Ala Ala Ser Leu Tyr Tyr Pro 50 S

Gln Gln Ala Ile Val Phe Ala Gln Val Ser Gly Glu Arg Leu Tyr Asp 65 70 80

Trp Gly Leu Arg Gly Tyr Ile Val Ile Glu Asp Leu Trp Lys Glu Asn 90 85 Phe Gln Lys Pro Gly Asn Val Lys Asn Ser Pro Gly Thr Lys Xaa 110

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(2) INFORMATION FOR SEQ ID NO: 334:

(A) LENGTH: 106 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS

SEQUENCE DESCRIPTION: SEQ ID NO: 334: ž

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Met Ala Pro Ser Leu Leu Leu Ala Pro Leu Cys Ser Leu Glu Ala Val Leu Ser

Ser Pro Leu Glu Lys Gln Cys Gln Leu Pro Gly Ile Phe 20 30 Ser Ala Gln Leu Leu ž Pro Leu Leu L 40 Cys Gln Leu Gln Leu Pro Cys

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Lys Gly Ile Val Xaa Pro Arg Cys Pro Ala Ser Leu Pro Gln Pro Pro 50 Leu Pro Leu His Cys Thr Glu Arg Xaa 75 Trp His 70 Pro Ala Pro Ser His 65

20

Pro His His Leu Pro Leu Gln Gly Gly Ser Ser Asn Met Glu Glu Xaa 85

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Asn Tyr Arg Gly Tyr Xaa Asp Ala Gln Leu 100

3

(2) INFORMATION FOR SEQ ID NO: 335:

35

(A) LENGTH: 50 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 335:

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Met Thr Thr Cys Leu Phe Gly Leu Leu Ser Cys Glu Met Ser Ala Gln 1 5

Val Ser Gln Lys Ser Cys Val Tyr Asp Glu Ser Glu Cys Phe Ser Ser 20 45

Val Gly Gln Leu Leu Ala Leu Leu Ile Leu Val Tyr Val Leu Pro Ser 15

Ile Xaa S0

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(2) INFORMATION FOR SEQ ID NO: 336:

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(A) LENGTH: 48 amino acids (B) TYPE: amino acid (i) SEQUENCE CHARACTERISTICS:

Met Thr Phe Ser Ser Leu Lys Leu Phe Val Leu Thr Cys Ile Ile Lys 1  $\phantom{0}$  15  $\phantom{0}$ Gln Arg Ser Leu Ser Ser Asn Leu Val His Val Leu Leu Gln Pro Ala 15 40 45 Gly Leu Glu Arg Phe Ile Ile Leu Arg Glu Val Cys Asn Glu Glu Ile  $20 \ \ 30$ Phe Lya Asp Val Leu Val Thr Glu Ile Ile Cys Leu Cys Met Cys 50 . 55 60 X. SEQUENCE DESCRIPTION: SEQ ID NO: 338: (A) LENGTH: 76 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

6 ઝ 30 25 Met Leu Ile Pro Leu Gin Cys Leu Phe Ser Ser Asp Arg Met Leu Thr 1 5 Phe Leu Thr Fro Trp Gln Lys Gly Glu Lys Cys Val Leu Gly Trp Val 20 25 (2) INFORMATION FOR SEQ ID NO: 338: Ε¥ (i) SEQUENCE CHARACTERISTICS: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 337: Phe Leu Ser Glu Ile Ser Xaa 35 40 (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 41 amino acids

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(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 337:

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Txp Lys Ser Ile Pro Gly Pro Ser Pro Arg Asn Gln His Arg 35  $40\,$ 

Leu Ala Leu Val Glu Ile Lys Leu Glu Asp Leu Gln Ser Gln Leu His 20 25

Met Leu Trp Lys Cys Ser Gln Asn Ile Ala Arg Cys Leu Leu Leu Leu 1 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 336:

(D) TOPOLOGY: linear

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Leu Tyr Ser Ile Lys Tyr Met Pro Pro Gln Lys Lys 65 70 75

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Met Arg Leu Phe Phe Ile Gly Phe Leu Leu Leu Phe Ser Phe Gly Leu  $10 \ 15$ 

SEQUENCE DESCRIPTION: SEQ ID NO: 342: (D) TOPOLOGY: linear

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Thr Glu 25 Ę Asp Leu Ala Lys 20 Arg Ser Ser

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INFORMATION FOR SEQ ID NO: 343: 3 으 SEQUENCE CHARACTERISTICS:

(A) LENGTH: 157 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 343: ž.

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Phe Ser Ser Leu Glu Phe Tyr Gln Lys Lys Lys Ser Arg Trp Pro

Asp Glu Cys Ile Pro Trp Glu Val Trp Thr Val Lys Val His Val Val 20 20 8

Ala Leu Ala Thr Glu Glu Glu Arg Gln Ile Cys Arg Glu Lys Val Gly
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Glu Lys Leu Cys Glu Lys Ile Ile Asn Ile Val Glu Val Met Asn Arg 50

His Glu Tyr Leu Pro Lys Met Pro Thr Gln Ser Glu Val Asp Asn Val 65 8

Phe Asp Thr Gly Leu Arg Asp Val Gln Pro Tyr Leu Tyr Lys Ile Ser 95

Phe Gln Ile Thr Asp Ala Leu Gly Thr Ser Val Thr Thr Thr Met Arg 100 33

Arg Leu lle Lys Asp Thr Leu Pro Ser Glu Arg Arg Trp lle Ser Gly 115 <del>6</del> Ser Leu Met Ala Pro Arg Pro Trp Leu Leu Gly Ile Ala Leu Leu 130 Ser

Leu Trp Ala Leu Glu Pro Ala Leu Gly His Trp Xaa 150 G1y 145 5

INFORMATION FOR SEQ ID NO: 344: 3 S

(A) LENGTH: 520 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 344: 3

25

Leu Pro Leu Pro Ala Ala Gly Arg Val Val Val 5 Phe Leu Met

Leu Ala Val Arg Arg Phe Gly Ser Arg Ser Leu Ser Thr Ala Asp Met 8

Leu Lys Ala 80 Glu Lys Glu Asp Asp Asp Lys Leu Leu Ala 60 Leu His Gln Asp Phe Pro Ser Val 90 95 8 Pro Pro Thr Lys Gly Leu Val Leu Gly Ile Tyr Ser Lys Ser Ala Gly Glu Asn Phe 55 Thr Leu Asn Ile Ser Gly 70 75 22 Thr Phe Tyr Gly 85 Val Pro Gln Phe Thr Gly Lys Leu Arg Glu Gly Lys Thr Arg 2

Leu Val Gly Leu Gly Lys Lys Ala Ala Gly Ile Asp Glu Glu Glu 100 100 Asn Trp His Glu Gly Lys Glu Asn Ile Arg Ala Ala Val Ala Ala Gly 125 Val 15 ನ

Gly Asp Ala Gin Ala Ala Glu Gly Ala Val Leu Gly Leu Tyr 150 Ser Val Glu Val Asp 140 Gln Ile Gln Asp Leu Glu 135 Arg 130 143 25

Pro

Ser

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Glu Tyr Asp Asp Leu Lys Gln Lys Lys Lys Met Ala Val Ser Ala Lys 170

Leu Tyr Gly Ser Gly Asp Gln Glu Ala Trp Gln Lys Gly Val Leu Phe 180 Gly Gln Asn Leu Ala Arg Gln Leu Met Glu Thr Pro Ala Asn 195 Ser Ala 3

11e 240 Met Thr Pro Thr Arg Phe Ala Glu Ile Ile Glu Lys Asn Leu Lys 210 Thr Glu Val His Ile Arg Pro Lys Ser Trp 230 235 Ser Ser Lys Ala gr Ser 225

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Glu Glu Gln Ala Met Gly Ser Phe Leu Ser Val Ala Lys Gly Ser Asp 255 8

Glu

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Pro Pro Val Phe Leu Glu Ile His Tyr Lys Gly Ser Pro Asn Ala 260 270 Ser Asn.Glu Pro Pro Leu Val Phe Val Gly Lys Gly Ile Thr Phe Asp 275 20

Leu Met Arg Ala Met Gly Gly Ala Ala Thr Ile Cys Ser Ala Ile Val Ser Ala Ala 320 Gly Ile Ser Ile Lys Ala Ser Ala Asn Met Asp 290 Asp 305 g 55

Pro Leu Cys Glu 335 Asn Met Pro Ser Gly Lys Ala Asn Lys Pro Gly Asp Val Val Arg Ala Asn Leu Pro Ile Asn Ile Ile Gly Leu Ala 325 330

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345 350

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Lys Asn Gly Lys Thr Ile Gln Val Asp Asn Thr Asp Ala Glu Gly Arg 355

퉏 Ile Leu Ala Asp Ala Leu Cys Tyr Ala His Thr Phe Asn Pro Lys 370 380

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5 Xaa Ile Leu Asn Ala Ala Thr Leu Thr Gly Ala Met Asp Val Ala Leu 385 390 395

Gly Ser Gly Ala Thr Gly Val Phe Thr Asn Ser Ser Trp Leu Trp Asn 415

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Ala Ala Val Leu Gin Ser Asp Thr Met Asp His Tyr Arg Thr Phe His 50 60

Val Ala Leu Arg Val Arg Ser Gly Ile Leu Glu Gln Thr Gly Ala Thr \$35\$

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Glu Asp Val Tyr Arg Leu Trp Leu Asp Gly Tyr Ser Val Thr Asp Ala 20

Met Thr Ser Glu Leu Asp Ile Phe Val Gly Asn Thr Thr Leu Ile Asp 1 .15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 346:

(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Lys Leu Phe Glu Ala Ser Ile Glu Thr Gly Asp Arg Val Trp Arg Met 420 425

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Pro Leu Phe Glu His Tyr Thr Arg Gln Val Val Asp Cys Gln Leu Ala 415 440

20 Asp Val Asn Asn Ile Gly Lys Tyr Arg Ser Ala Gly Ala Cys Thr Ala 450 455

Ala Ala Phe Leu Lys Glu Phe Val Thr His Pro Lys Trp Ala His Leu 475 470 475

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Ser Lys Gly Thr Lys Lys Asp 115

Leu Asp Asp Ile Ser Thr Lys Thr Gly 120 125

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Ile Thr Leu Lys Ser Cys Arg Arg Gln Phe Asp Asn Pho Lys Arg Val 130 140

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Ile Phe Gln Ile Pro Pro Ser Arg Gln Ala Leu Leu Ile Glu Arg Tyr 85 90 95

Met Leu Glu Arg Leu Leu His Ala Pro Pro Lys Leu Leu His Gln Leu 65 70 75 80

Tyr Ala Phe Asp Glu Ala Phe Val Arg Glu Val Leu Gly Lys Lys Leu 100

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Asp Ile Ala Gly Val Met Thr Asn Lys Asp Glu Val Pro Tyr Leu Arg 495

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SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 345:

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SEQUENCE DESCRIPTION: SEQ ID NO: 345:

(D) TOPOLOGY: linear TYPE: amino acid LENGTH: 39 amino acids

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Gln Ser Leu Arg Leu Asn Ala ,35

Gly Lys Asp Ser Ile Asp Ile Asp Ile Ser Ser Arg Arg Arg Glu Asp 25

Thr Ile Leu Phe Leu Phe Leu Gln Leu Ser Ala Leu Arg Leu Ile Val 1 5

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INFORMATION FOR SEQ ID NO: 346:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 234-amino acids

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Phe Ser Gln Asp Asn Ala Xaa 515

Lys Gly Met Thr Gly Arg Pro Thr Arg Thr Leu Ile Glu Phe Leu Leu 500 510

Gln His Phe Leu Leu Ser Asp Arg Leu Ala Arg Asp Tyr Ala Ala Ile 165 170 175 Phe Lys Val Val Glu Glu Wet Arg Gly Ser Leu Val Asp Asn Ile Gln 145 150 150

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Val Phe Phe Ala Asn Asn Arg Phe Glu Thr Gly Lys Lys Lys Leu Gln 180 185

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Tyr Leu Ser Phe Gly Asp Phe Ala Phe Cys Ala Glu Leu Met Ile Gln 195 200 205

Asp Leu Asp Arg Asn Phe Ser Arg Thr Xaa 225 230 Trp Thr Leu Gly Pro Val Asp Ser Gln Met Asp Map Met Asp Met 210 215

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Asn

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(2) INFORMATION FOR SEQ ID NO: 347:

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Ξ SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 169 amino acids

SEQUENCE DESCRIPTION: SEQ ID NO: 347:

Met Ala Ala Ala Val Ala Gly Met Leu Arg Gly Gly Leu Leu Pro Gln

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Ala Gly Arg Leu Pro Thr Leu Gln. Thr Val Arg Tyr Gly Ser Lys Ala 20

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Val Thr Arg His Arg Arg Val Met His Phe Gln Arg Gln Lys Leu Met 35

Ala Val Thr Glu Tyr Ile Pro Pro Lys Pro Ala Ile His Pro Ser Cys 50 2 Leu Pro Ser Pro Pro Pro Gln Glu Glu Ile Gly Leu Ile Arg 65 70 78 80

Leu Leu Arg Arg Glu Ile Ala Ala Val Phe Gln Asp Asn Arg Met Ile 85 Cys Gln Asn Val Ala Leu Ser Ala Glu Asp Lys Leu Leu Ile 100 Ala Val 15

Pro Ala Ala Glu Thr Gln Asp Pro Asp Glu Gly Leu Pro Gln 115 뵱 Ala

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Cys Pro Phe Leu Trp Gly Thr Thr Cys Cys Trp Ser Val Lys Ser Pro 145 Gly Pro Glu Ser Pro Ser Trp Arg Ile Pro Ser Thr Lys Ile Cys 130 Pro

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Arg Ser Arg Arg Trp Tyr Gly Ser Xaa 165 ജ

INFORMATION FOR SEQ ID NO: 348: 3 35

(A) LENGTH: 43 amino acids (1) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 348:

Lys Arg Ser Phe Leu Leu Pro Leu Leu Val Gly Phe Leu Asp 10 15 껿 5

Thr Ala His Leu Ile Leu Leu Glu Thr Leu Ser Val Cys Leu Trp Leu 30

Pro Ser Leu Ile Asp Ser Arg Cys Val Met Ser 35 40 S

INFORMATION FOR SEQ ID NO: 349: 3 55

(A) LENGTH: 78 amino acids (1) SEQUENCE CHARACTERISTICS: TYPE: amino acid

(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 349: ž

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Met Lys Glu Gly Pro Pro Cys Lys Arg His His Tyr Tyr Gln Asn Cys

Phe Gly Glu Thr Asn Gln Ile His 25 Gly Ala Lys Leu Leu Val Ser Leu 20 Ś

Leu Leu Glu Thr Gln Val Gly Thr Glu Lys Gly Glu Arg Ile Trp 35

Glu Glu Lys Trp Arg Ile Ser Ser Thr Val Leu Phe Ile Ser Val Asn 2

Ser Tyr Val Glu Gly Ser Val Leu Glu Ile Lys Leu Phe Tyr 65  $^{75}\,$ 2

(2) INFORMATION FOR SEQ ID NO: 350:

2

SEQUENCE DESCRIPTION: SEQ ID NO: 350: (X

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Met Ser Glu Ile Leu Ser Leu Leu Phe Cys Leu Leu Gly Pro Ala Leu 10 15 15

Asp Glu Arg Arg Glu Glu Lys Asp 20

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(2) INFORMATION FOR SEQ ID NO: 351:

35

(A) LENGTH: 274 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 351:

6

Met Ser Ser Ala Gly Thr Ala Thr Pro Leu Glu Met Asp His Lys Leu 1 15 15

5

Thr Ser Gln Pro Gly Arg Pro Ser Phe Tyr Cys Asn Ser Arg His Ser 20

lle Val Gly Ser Ser His Gln Leu Gly Phe Trp Phe Ser His Leu Glu 35

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Ser Ser Gly Leu Lys Val Phe Gln Val Ser Leu Pro Cys Glu Cys Val 50 60 Asn Leu Pro Thr Arg Ile Ala Ser Val Val Leu Ser Leu Met Ser Leu 65 70 70 70 55

Leu Val Val Gly Oln Ala Pro Ala Trp Glu Gly Ser Leu Leu Arg Gly 85 99 S

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Arg Pro Ala Gly Gly Ala His Leu Cys Ala Met Xaa Val Ile Glu Gly 100 105

S Gly Gln Val Ser Gln Val Leu Pro Ala Leu Ser Leu Gly Leu Val Phe 130 135

5 Leu Cys Gln Gly Thr Val Glu Lys Val Ser Gly Ala Ala His Cys Ser 145

5 Thr Xaa Arg Cys Ser Arg Pro Tyr Phe Ser Ser His Lys Gly Val Ala 180

20 Ala Thr Leu Ala Leu Thr Cys His Cys Asp Lys Val His Val Ala Gly
195 200

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Val Thr Gln Ala Lys Trp Asn Ser Trp Pro Ser Arg Arg Asn Ala Gly
35 40 45

Glu Ala Xaa Val Arg Ser Ser Lys Lys Trp Ile Pro Lys Ala Leu Xaa 20 25 30

Met Ser Ile Ser Gly Thr Asp Gly Leu Ile Leu Leu Leu Val Gly Leu 1 5 10

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SEQUENCE DESCRIPTION: SEQ ID NO: 354:

(A) LENGTH: 52 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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35 Ile Xaa

(2) INFORMATION FOR SEQ ID NO: 352:

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ε SEQUENCE CHARACTERISTICS:
(A) LENGTH: 47 amino acids
(B) TYPE: amino acid

Ě SEQUENCE DESCRIPTION: SEQ ID NO: 352: (D) TOPOLOGY: linear

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Met Ile Phe Thr Ser Val Thr Lys Gly Ile Leu Leu Ile Ala Leu Trp 1 5

50 Val Pro Leu Phe His Phe Met Leu Ile Asp Ser Ile Leu Gly Pro Ser  $20 \ \ 25$ Arg Leu Leu Thr Asp Gly Val Pro Phe Asn Pro Trp His Val Xaa 35

(2) INFORMATION FOR SEQ ID NO: 353:

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(i) SEQUENCE CHARACTERISTICS:

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Leu Val Val Asp Val Gly Glu Arg Ile Leu His Gly Gln Arg Glu Val 115 120

Ser Leu Leu Cys Cys Leu Pro Trp Gln Cys Ser Gly Gly Gly Phe Pro 165 170 175

Leu Gly Lys Asp Txp Ala Ile Glu Gln Arg Arg Thr Cys Glu Ser 210 215

Asp Xaa Glu 2 225 XAA XAA Pro Phe Thr Leu Ala Gly Leu Val Leu Val Leu 230 235

Cys Gln Val Val Leu Val TTP Ile Pro Gln Leu Gly Asp Lys 250 255

Arg Gly Met Thr Arg Leu Gly Arg Val Ser Leu Thr Ser Ser 260 265

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Phe Ala Leu His 50

(2) INFORMATION FOR SEQ ID NO: 355:

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(i) SEQUENCE CHARACTERISTICS: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 355: (A) LENGTH: 132 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Glu His Cys Leu Tyr His Ser Val His Gly Ile Asn Pro Tyr Ile
1 19 15

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His Lys Asn Thr His Pro Ser Ile Asn Ile Tyr Met Val Trp Asp Glu 20 25 30

3

Gln Val Asn Ser Phe Glu Arg Glu Phe Val Pro Phe Phe Phe Leu Ile  $35 \hspace{1cm} 40 \hspace{1cm} 45$ 

Ile Leu Leu Asn Cys Cys Gln Leu Ser Asn Lys Gln Thr Glu Lys Leu  $50 \ \ 55$  60

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Arg Leu Asn Thr His Ile Leu Pro Val Phe Ser Tyr Ser Asp Ser Ile 90 95 Phe Gly Lys Thr Leu His Thr Pro Phe Leu Ser Ser Ala Leu Lys Tyr 65 70 75 80

S

Leu Thr Cys His Leu Ile Leu Ala Ser Tyr Phe Ser His Val Tyr Leu 100 105 110 100

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Met Lys Thr

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(2) INFORMATION FOR SEQ ID NO: 354:

(i) SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 353:

(A) LENGTH: 3 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

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Pro Val Thr Cys Ile Cys Tyr Leu Asn Arg Lys Lys Asn Ile Gln Lys 115

Lys Lys Asn Xaa 130

(2) INFORMATION FOR SEQ ID NO: 356: 9

(A) LENGTH: 204 amino acids (i) SEQUENCE CHARACTERISTICS:

TYPE: amino acid

15

SEQUENCE DESCRIPTION: SEQ ID NO: 356: Ŕ

Gly Ser Arg Asp His Leu Phe Lys Val Leu Val Val Gly Asp Ala 5 Met

Phe Asp Ser Gln Asp Tyr Ser Gln 25 Leu Val Ser 뀵 Ala Val Gly

2

Thr Val Gly Val Asp Phe Ala Leu Lys Val 40 Lys His Tyr Lys Ser 35 Ser 25

Trp Asp Leu Gln Leu Glu Ile Val Arg ጟ Ş Gln Trp Ser

Leu Tyr Tyr Arg Asp 80 Ala Gly Gln Glu Arg Phe Thr Ser Met Thr Arg 65 75 75 8

Thr Thr Phe Phe Asp Val Thr Asn Ala 90 Ala Ser Ala Cys Val Ile Met

Lys Leu Thr Leu 110 Ser Asn Ser Gln Arg Trp Lys Gln Asp Leu Asp 105 Ser

35

Pro Asn Gly Glu Pro Val Pro Cys Leu Leu Leu Ala Asn Lys Cys Asp 125 6

Glu Asn Gly Phe Thr Gly Trp Thr Glu Thr Ser Val Lys Glu Asn Lys 145 5

Ser Pro Trp Ala Val Ser Arg Asp Gin Ile Asp Arg Phe Ser Lys 130

Se

Asn Ile Asn Glu Ala Met Arg Val Leu Ile Glu Lys Met Met Arg Asn 175 20

Glu Asp Ile Met Ser Leu Ser Thr Gln Gly Asp Tyr Ile Asn 180 180 걡 Ser

Trp Ser Cys Cys Xaa 200 Leu Gln Thr Lys Ser Ser Ser 195 55

(2) INFORMATION FOR SEQ ID NO: 357:

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(A) LENCTH: 47 amino acids SEQUENCE CHARACTERISTICS: 3

(B) TYPE: amino acid (D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 357: Ŧ

Met 11e Ser Leu I1e Phe Gln Leu Glu Glu Glu Lys Leu Val Glu Lys  $1 \ \ \, 1$ 

Leu Phe Phe Leu Lys Lys Gly Ser Gln Gly Ser Leu Arg Gly Val Pro Arg His Met Arg Val Lys Ile Val Phe 8 Phe Asn Leu Phe Phe 9

(2) INFORMATION FOR SEQ ID NO: 358:

12

(A) LENGTH: 73 amino acids (i) SEQUENCE CHARACTERISTICS:

2

TYPE: amino acid

SEQUENCE DESCRIPTION: SEQ ID NO: 358: ž

53

Met Thr Tyr Val Thr Cys Leu His Val Cys Leu Leu Val Glu Phe Leu

Phe Trp Phe Thr Gly Leu Arg Gly Phe Ser Glu Tyr Leu Trp Pro Gln 15 Asn Ser Gln Leu Thx Asn His Arg Lys Tyr Tyr Phe Leu Ser Tyr Gly 20 25

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Gin His Thr Ser Phe His Pro Asn Arg Asn Glu Ile Asn Phe Val Ser S0 60 35

Thr Asp Asn Arg Ile Trp Val Thr Xaa 65

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(2) INFORMATION FOR SEQ ID NO: 359:

(A) LENGTH: 102 amino acids (i) SEQUENCE CHARACTERISTICS

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(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 359:

20

Lys Glu Gly Glu Tyr Ile Lys Leu Lys Val Ile Gly Gln Asp Ser Ser 20 20 20

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Ľys Glu Ile His Phe Lys Val Lys Met Thr Thr His Leu Lys Lys Leu 15 45 Glu Ser Tyr Cys Gln Arg Gln Gly Val Pro Met Asn Ser Leu Arg Phe

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Gly His Ser Thr Val Xaa 100 Gly Met Glu Glu Glu Asp Val Ile Glu Val Tyr Gln Glu Gln Thr Gly 95 Gln Val Leu Glu Ala Pro Gly Val Tyr Val Phe Gly Glu Leu Leu Asp \$45\$Met Ser Ala Glu Val Lys Val Thr Gly Gln Asn Gln Glu Gln Phe Leu 1 15 Leu Leu Val Gly Leu Val Tyr Leu Val Ser His Leu Ser Gln Arg
35 40 45 Leu Phe Glu Gly Gln Arg Ile Ala Asp Asn His Thr Pro Lys Glu Leu 65 70 75 80 Met Pro Așn Val Arg Glu Leu Ala Glu Ser Asp Phe Ala Ser Thr Phe 50 55 (2) INFORMATION FOR SEQ ID NO: 360: Glu Ala Arg Asn Leu Pro Pro Leu Thr Glu Ala Gln Lys Asn Lys Leu 95 (2) INFORMATION FOR SEQ ID NO: 361: Leu Ala Lys Ser Ala Lys Gly Ala Ala Leu Ala Thr Leu Ile His . \$20\$SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: Pro Gln Trp His Leu Gly Asn His Ala Val Glu Pro Val 5 10 15 Thr Val Phe Ala Tyr Gly Thr Tyr Ala Asp Tyr Leu Ala 70  $\,$  75  $\,$  80  $\,$ SEQUENCE DESCRIPTION: SEQ ID NO: 361: Leu Leu Phe Leu Leu Met Met Leu Gly Val Arg Gly 20 25 30 SEQUENCE DESCRIPTION: SEQ ID NO: 360: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 179 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear LENGTH: 48 amino acids 5 5 \$ 6 3မ 25 20 8 55 S Met Lys Ser Ser Ser Leu Phe Phe Phe Phe Leu Ala His Phe Ile His 1 15 2

568

Ser His Asp Leu Pro Gly Leu Cys Arg 20 25

(2) INFORMATION FOR SEQ ID NO: 363:

Ξ SEQUENCE CHARACTERISTICS acids

ž SEQUENCE DESCRIPTION: SEQ ID NO: 363: (A) LENGTH: 224 amino e (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Lys Phe Ala Ala Ser Cly Xaa Phe Leu His His Met Ala Cly Leu 1 15

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Arg 65

Leu Leu

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Ser Ser Lys Leu Ser Met Ser Lys Ala Leu Pro Leu Thr Lys Val 20 25 30

Val Gln Asn Asp 35 Ala Tyr Thr Ala Pro Ala Leu Pro Ser Ser Ile Arg 40 45

쿢 Pro Pro Lys Glu Leu Pro Ala Ala Glu Pro Val Leu Ser Pro Leu Glu 65 70 80 Lys Ala Leu Thr Asn Met Ser Arg Thr Leu Val Asn Lys Glu Glu 50 55

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Met Gly Phe

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Thr Ser

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INFORMATION FOR SEQ ID NO: 362:

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 362: (A) LENGTH: 25 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

(i) SEQUENCE CHARACTERISTICS:

Asn His Xaa

Asp Ile Gln Arg Gln Asp Leu Ser Ala Ile Ala Arg Thr Leu Xaa Lys 165 170 Leu Asp Gln Arg Asn Gln Arg Leu Glu Val Asp Tyr Ser Ile Gly Arg 145 150

Glu Asp Leu Val Ile Glu Ala Val Tyr Ala Asp Val Leu Arg Gly Ser 130  $$125\,$ 

Tyr Ala Val Leu Leu Glu Ala Leu Ala Leu Arg Asn Val Arg Gln Leu 115 120 125 Arg His Leu Ser Val Val Thr Leu Ala Ala Lys Val Lys Cys Ile Pro 100 105 110

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Ser 160

Ser

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Leu His 255 Met Lys Thr 175 Phe Pro Ser 270 Leu Gln Glu Leu Leu Ser Lys Cys Arg Thr Cys. 120 Glu Thr Arg Glu Val Leu Thr Pro Thr Ser Thr Ser Asp Asn Glu Thr 190 Ser Asn Ile Arg Ser Gln His Ala Glu Glu Gln 230 240 Leu Arg Ser Cys 285 Ser Gln Asp Pro Glu Val Ala Leu Ser Leu 295 Phe Ser His Met Gln Gln His Asp 310 Ile Glu Ser Thr Ile His Val 330 335 Leu His Gln Asp His Ile Thr Phe Ala Met Leu Leu Ala Arg Ile 5 Lys Leu Lys Gly Thr Val Gly Glu Pro Thr Tyr Asp Ala Glu Phe Gln Leu Gln Glu Glu Ala Lys Glu Arg 135 Pro Glu Asn Ser Ser Val Lys Glu Tyr Arg Met Glu Val Pro Ser 215 Lys Asp Asp Glu Gly Ala Thr Pro Ile Lys Arg Arg Val 150 Leu Pro Ser Asn Asn Gly Arg Tyr Asp Asp Cys Lys Glu Phe Lys Asp 250Ser Cys Ile Ser Asp 170 Ser Ser Ile Ile Asp Pro Gly Thr Glu Gln Asp 195 200 205 Glu Glu Glu Ser Glu 265 Gln Ala Ala Ser Xaa 345 SEQUENCE DESCRIPTION: SEQ ID NO: 365: Asp Ser Asp Leu Ala 280 (A) LENGTH: 467 amino acids (i) SEQUENCE CHARACTERISTICS: TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 365: Ser Arg Gly Leu 310 Cys Arg Thr Ser Asp Glu Glu His Thr Val Asp 165 Ę Ser Thr Leu Ala Leu Thr Arg Ile Ser Gly Lys Gly 340 Ser Gly Gln Ala Leu Pro 290 Thr Ser Ile Ser Ala Val 275 Leu 325 Ala Leu Ile Pro Thr Asn Aet Gln Gln Arg 1 130 Ser Cys Gly His 305 Cys Ser Lys Asp 260 Ser Glu Asp 돮 Ile Leu Asp (X Arg Asp Phe 225 ž 14 S βg Met S 2 8 45 8 2 22 8 35 6 တ္တ 55 Lys Glu 175 Ser Asn 80 Leu Arg 95 Phe Ala 15 Asp 160 Leu Leu Sar Val His Thr Pro Lys Gln Leu Asn Pro 110 Gly Thr Lys Met Thr Val Asn Asn Leu His Pro Arg Val Thr Glu Glu 99 Asp Lys Tyr Asn Asn Arg Cys Leu Asp Gly Gln 135 Phe Lys Ile Lys Leu Xaa 220 Phe Ser Glu Ala Asn Cys Ala Asn Leu Ile Ser Thr Leu Ile 55 Ala Arg Pro Pro Ile Leu Lys Ala Leu Phe Lys Ser Ser 200 Met Asp Glu Arg Thr Phe Leu Asn Asn 25 His Phe Leu Leu Lys Val Gln Ser 45 Phe Val Lys Lys Asp. 125 Ser Lys Arg Ser Pro Ser Met Lys 170 Met Ser Lys Asn Cys Ile Lys Leu Leu Cys Glu Asp Pro Val 1 5 Phe 큪 Asn 190 Arg Val Glu Ile Ser Lys Ala Ser Ala Ser Leu Asn Gly Asp 90 Met Asn Gly Asn Val Ile 155 Gln Tyr Gln Asn Leu Gln Ser Asp 70 Ser Ser (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 364: Val Cys Gly Ala Leu 105 Ala Ser a Glu Val Val F 120 Gln Pro Thr Glu | 215 Asn Ser 185 Gin Pro Ile Leu Leu Arg Leu Ser Asp 165 SEQUENCE CHARACTERISTICS TYPE: amino acid INFORMATION FOR SEQ ID NO: 364: (D) TOPOLOGY: linear Thr e Phe Cys НÌВ Glu Val Asp Pro Asp Thr 195 Glu Tyr Ile Lys Cys Ile Leu 20 Жet Val Ala Leu Pro Arg Arg Val 15 E Ly8 녎 Phe Asp Ile Val Glu Leu 100 His Pro Gly Ala Ile Thr Ala Tyr 130 Pro Met Lys Cys Asn Ser Xaa Thr Asn ile Val Tyr Thr 35 Asn Leu Ile Ser 3 8 180 Ala Leu Ala 3 Ala 210 Gln val Сħп Leu Val Ser ਫ਼ੋ 를 S 3

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6 25 20 5 45 35 30 2 55 50 Glu Gln Phe Gly Ile Trp Leu Asp Ser Ser Ser Pro Glu Gln Thr Val 85 90 95 Ser Cys Leu Pro Ala Phe Lys Asp Leu Ile Ala Lys Val Gln Ala Asp 65 70 75Arg Ile Gln Gly Leu Thr Val Glu Gln Ala Glu Ala Val Val Arg Leu  $50 \ \ 60$ His Phe Leu Arg Gly Asn Glu Ile Val Leu Ser Ala Gly Ser Thr Pro 35 40 Ala Ser Gly His Val Glu Asp Leu Ala Ala Glu Gln Asn Thr Gln Ile 180 Ala Met Ala His Met Phe Val Ser Thr Asn Leu Gly Glu Ser Phe Met 130  $$13^\circ$$ Ile His Arg Leu Leu Leu Ile Gln Ala Phe Arg Pro Asp Arg Leu Leu 115 120 128 Val His Leu Ala Pro Gly Trp Leu Met Gln Leu Glu Lys Lys Leu His 225 230 230 235 Ala Ile Asn Thr Ala Val Lys Ser Gly Arg Trp Val Met Leu Lys Asn 210 225 Thr Ser lle Ala Ile Gly Ser Ala Glu Gly Phe Asn Gln Ala Asp Lys 195 200 205 Val Lys Pro Asn Thr Pro Val Leu Met Cys Ser Val Pro Gly Tyr Asp 165 170 Ser Ile Met Glu Gln Pro Leu Asp Leu Thr His Ile Val Xaa Thr Glu 145 150 150 Pro Tyr Leu Leu Tyr bhe Leu Leu Ala Trp Phe His Ala Ile Ile Gln Glu Arg Leu 305 310 Ser Ile Pro Val Ser Arg Ile Cys Lys Ser Pro Asn Glu Arg Ala Arg 290 295 Phe Glu Pro Pro Pro Gly Xaa Lys Ala Asn Met Leu Arg Thr Phe Ser 275 280 285 Asn Pro Lys Val Pro Val Asn Leu Leu Arg Ala Gly Arg Ile Phe Val 260 265Ser Leu Gln Pro His Ala Cys Phe Arg Leu Phe Leu Thr Met Glu Ile 245 250 Asp Leu Arg Ser Xaa Cys Asp Thr Val Asp Thr Trp Leu Asp Asp Thr Arg Tyr Ala Pro Leu Gly Trp Ser Lys Lys Tyr Glu Phe Gly Glu Ser 325 330 335 Trp Ser Glu Glu Thr Pro Ala Thr Pro Ile Gly Gln Ala 100 105 20 25

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(2) INFORMATION FOR SEQ ID NO: 366:

(1) SEQUENCE CHARACTERISTICS

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Lys Thr Asn Ala Gly Pro Arg Asp Arg Glu Leu Trp Val Gln Arg Leu  $20 \ 30$ 

Met Ala Asp Glu Ala Thr Arg Arg Val Val Ser Glu Ile Pro Val Leu 1 5 10

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 366:

Lys Glu Glu Tyr Gln Ser Leu Ile Arg Tyr Val Glu Asn Asn Lys Asn 45

45

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Trp Phe Gly Lys Cys Trp Tyr Ile His Asp Leu Leu Lys Tyr Glu Phe 65 70 75 80

Ala Asp Asn Asp Trp Phe Arg Leu Glu Ser Asn Lys Glu Gly Thr Arg 50 55

SS

Ile Ala Val Pro Glu Leu 100

Asp Gly Lys Thr Ala Lys Met Tyr Arg Gly 105 110

Asp Ile Glu Phe Asp Ile Pro Ile Thr 85

Tyr Pro Thr Thr Ala Pro Glu 90 95

25

His Asp Gln 465

20

Pro Ala Gln Gln Arg Arg Glu Ser Pro Pro Tyr His Thr Gly Cys Gly  $450 \hspace{0.25cm} 460 \hspace{0.25cm}$ 

Ala Val Gly Gly Val Ala Pro Arg His Pro Asp Ala Leu Leu Ala Gly 435 440 445

<u></u> 12 Glu Phe Asp Gln Arg Leu Leu Asn Thr Phe Leu Glu Arg Leu Phe Thr 385 390 395 Ala Lys Gly Arg Gln Asn Ile Ser Pro Asp Lys Ile Pro Trp Ser Ala 355 360 365 His Lys Asp Ile Gln Met Pro Asp Gly Met Gln Ala Arg Gly Val Cys \$420\$Thr Arg Ser Phe Asp Ser Glu Phe Lys Leu Ala Cys Lys Val Asp Gly
415 Leu Lys Thr Leu Met Ala Gin Ser Ile Tyr Gly Gly Arg Val Asp Asn 370 375

572

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Gly Lys Ile Cys Leu Thr Asp His Phe Lys Pro Leu Trp Gly Gln Glu 115 120 126

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His Gly Ser Gly Ala Gly Ser 140 Ser Cys Ala Gln Ile Trp Thr Ser Ser 130

7 Asn Pro Xaa 150 Met Xea Gly Ser Gly 145

INFORMATION FOR SEQ ID NO: 367 3 2

(A) LENGTH: 373 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 367 (x1)

2

Tyr Asp Gly Thr Lys Glu Val Pro Met Asn Pro Val Lys Ile Tyr 5

Gin Val Cys Asp Ile Pro Gin Pro Gin Gly Ser Ile Ile Asn Pro Gly 20 20

2

Ser Thr Gly Ser Ala Pro Trp Asp Glu Lys Asp Asn Asp Val Asp Glu 35

25

Glu Asp Glu Glu Asp Glu Leu Asp Gln Ser Gln His His Val Pro Ile 50 60

Phe Leu Asn Ile Asn Gly Ser Pro Met Ala Pro 70 80 Gln Asp Thr Phe Pro 65 2

Ala Ser Val Gly Aen Cys Ser Val Gly Asn Cys Ser Pro Glu Ala Val 85

Trp Pro Lys Thr Glu Pro Leu Glu Met Glu Val Pro Gln Ala Pro Ile 100 35

Gln Pro

Asp Leu Asp Ile Lys Phe Gln fyr Arg Gly Lys Glu fyr Gly Gln 130 Phe Tyr Ser Ser Pro Glu Leu Trp Ile Ser Ser Leu Pro Met 115 돸 6

Asn Pro Gln Gly Cys Arg Leu Phe Tyr Gly Asp 150 150 Ser Met Thr Val 14 5

Pro Asp Gln Glu Glu Leu Phe Gly Pro Val Xaa Leu 165 176 Leu Gly Pro Met 45

Lys Leu Leu Asp Val Met Asp Arg Gly Leu Ile Asn Glu Lys Gln Glu Gln Val Lys Phe Pro Gly Pro Glu His Ile Thr Phe Thr Ser 1 Lys Leu S

Leu Glu Val Ser Gly His Ala Ile Tyr Ala Ile Arg Leu Cys Gln Cys 210 210

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Ser Leu Val Ala Pro Asn 235 Lys Val Tyr Trp Ser Gly Pro Cys Ala Pro 225 8

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Leu Ile Glu Arg Gln Lys Lys Val Lys Leu Phe Cys Leu Glu Thr Phe. 250

Leu Ser Asp Leu lle Ala His Gln Lys Gly Gln Ile Glu Lys Gln Pro 260 265

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Glu ile Tyr Leu Cys Phe Gly Glu Glu Try Pro Asp Gly Lys 275 Pro Phe

Leu Glu Arg Lys Leu Ile Leu Val Gln Val Ile Pro Val Val Ala 290 Pro

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320. 320. Ser Val Arg Leu Gln Ile Ser Thr Pro Asp Ile Lys Asp Asn 335  $\mbox{Arg Met}$  lie Tyr Glu Met Phe Ser Gly  $\mbox{Asp}$  Phe Thr  $\mbox{Arg Ser}$  Phe 305 Ser Gly 2

ile Val Ala Gin Leu Lys Gin Leu Tyr Arg ile Leu Gin Thr Gin Glu 345 Ser Trp Gln Pro Met Gln Pro Thr Pro Ser Met Gln Leu Pro Pro Ala 360

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Pro Pro Gln Xaa 370 3

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(2) INFORMATION FOR SEQ ID NO: 368:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 368: SEQUENCE CHARACTERISTICS:
(A) LENGTH: 83 amino acids (D) TOPOLOGY: linear

35

Met Gly Ser Ser Val Leu Pro Phe Cys Val Cys Val Thr Ser Pro Ser 6 Leu Gly Gly Arg Cys Ile Gln Gly Arg Phe Ala Ser His Ser Lys Phe 20 20 Phe Gly Arg Lys Thr Ala Ser Phe Gly Ala Val Gly Glu Thr 35 45

Trp Gly

Pro Pro Asp Gln Glu Pro Gln Lys Glu Thr Glu Pro Ala Thr Ser Ser 50 60 S

His Ala Arg Pro Trp Ala Arg Val Ile Gly Leu Arg Ile Trp Pro Gln 65

Pro Asn Xaa

55

INFORMATION FOR SEQ ID NO: 369: 3

(i) SEQUENCE CHARACTERISTICS 575

(A) LENGTH: 21 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

(XX) SEQUENCE DESCRIPTION: SEQ ID NO: 369:

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Met Leu Leu Ser Val Ala Ile Phe Ile Leu Leu Thr Leu Val Tyr Ala 1 15

Tyr Trp Thr Met Xaa 20

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15 2

(D) TOPOLOGY: linear

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Leu Lys His Leu Ser Ser Gly Asp Leu Leu Arg Asp Asn Met Leu Arg 45

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Pro Gin Ala Glu Ala Leu Asp Arg Ala Tyr Gin Ile Asp Thr Val Ile 100 105

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SEQUENCE DESCRIPTION:

SEQ ID NO: 372:

(A) LENGTH: 51 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

25

Met Arg Ala Val Phe Pro Cys Cys Pro Phe Leu Thr Leu Met Leu Pro
1 5 10 15

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Leu Glu Cys Leu Val Gly Met Ile Met Cys Tyr Leu Gly Ile Ser 20 25 10

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**Lys Хаа Хаа** 

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Phe Thr Asp Thr Arg Lys Thr Ala Gly Leu Lys Lys Lys Lys Lys Lys Lys 45

35

(2) INFORMATION FOR SEQ ID NO: 372:

(i) SEQUENCE CHARACTERISTICS:

30

Lys Thr Ala Lys Gly Gly Gly Gln Glu Ala Leu Thr Cys Thr Xaa 65 70 75

Trp Ala Ile Lys Ala Gin Leu Lys Ile Glu Asn Lys Asp Leu Asp Asn 50 60

Lys Ile Ile Arg Cys Ser Ile Thr Ala Val Ser Asp Ser Ser Thr Ser 35 40 45 .

Leu Lys Ile Ser Met Gly Lys Tyr Leu Ser Ile Pro Thr Val Leu Leu 20 25

6

45 Asn Leu Asn Val Pro Phe Glu Val Ile Lys Gln Arg Leu Thr Ala Arg 115 120 125

Trp Ile His Pro Ala Ser Gly Arg Val Tyr Asn Ile Glu Phe Asn Pro . 130

50 Pro Lys Thr Val Gly Ile Asp Asp Leu Thr Gly Glu Pro Leu Ile Gln 145

Glu Asp Gln Arg Glu Asp Asp Lys Pro Glu Thr Val Ile Lys Arg Leu Lys Ala Tyr 165 170 175 Thr Lys Pro Val Leu Glu Tyr Tyr Gln Lys Lys Gly Val 180 185

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Leu Glu Thr Phe Ser Gly Thr Glu Thr Asn Lys Ile Trp Pro Tyr Val 195 200 205

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INFORMATION FOR SEQ ID NO: 370:

(i) SEQUENCE CHARACTERISTICS

X. SEQUENCE DESCRIPTION: SEQ ID NO: 370: (A) LENGTH: 227 amino acids (B) TYPE: amino acid

Met Gly Ala Ser Ala Arg Leu Leu Arg Ala Val Ile Met Gly Ala Pro 10

25

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Met Phe Leu Asn Cys Glu Ile Leu Glu Tyr Cys Tyr Tyr Leu Thr Gln 1 15

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<u>X</u>.

SEQUENCE DESCRIPTION: SEQ ID NO: 371:

(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 371:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 79 amino acids

S

Val Thr Pro 225

25

Gly Thr Glu Ile Gly Val Leu Ala Lys Ala Phe Ile Asp Gln Gly Lys 50 55 60

Leu Ile Pro Asp Asp Val Met Thr Arg Leu Ala Leu His Glu Leu Lys 65 70 75

35

Asn Leu Thr Gln Tyr Ser Trp Leu Leu Asp Gly Phe Pro Arg Thr Leu 95

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SS

INFORMATION FOR SEQ ID NO: 373:

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(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 61 amino acids

576

Tyr Ala Phe Leu Gin Thr Lys Val Pro Gin Arg Ser Gin Lys Ala Ser 210 215 220

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577

(B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 373: (x Phe Leu Met Arg Met His Leu Cys Phe Cys Lys Tyr Cys Cys Ser 10 F

Phe Ile Val Thr Pro Thr Ser Thr Ser Asn Thr Ala Ser Tyr Leu Trp 20

2

Ser Ser Trp Ala Cys Pro Trp Ile Ser Ala Ser Met Ala Gly Arg Gly 35

Glu Xaa Asn Ala Val Thr Arg Glu Gly Leu Pro 3 8 뀵

2

(2) INFORMATION FOR SEQ ID NO: 374:

2

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 40 amino acids

TYPE: amino acid

SEQUENCE DESCRIPTION: SEQ ID NO: 374: (D) TOPOLOGY: linear (Xi

25

Met Ser Leu Leu Asn Thr His Thr Leu Cys Phe Val Leu Phe Cys Phe 1 5 10

ile Asn Glu Glu Lys Leu Ala Asn His Leu Ala Phe Arg 20 30 Ser Ę 뀵 8

Ile Leu Phe Phe Ile Val Phe Xaa 35 40

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(2) INFORMATION FOR SEQ ID NO: 375:

(A) LENGTH: 44 amino acids (1) SEQUENCE CHARACTERISTICS: <del>송</del>

TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 375:

45

Met Cys Ser Gly Gln Ser Gln Val Trp Lys Met Ala Leu Gln Ala Leu

Asp Ser Glu Thr Val Val Ile Leu Pro Asp Met His Leu Ile Leu Ser 20 S

Leu Arg Leu Ile His Asn Ala Arg Pro Cys Leu Xaa 35

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(2) INFORMATION FOR SEQ ID NO: 376:

(A) LENGTH: 203 amino acids (1) SEQUENCE CHARACTERISTICS:

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578

(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 376:

Met Leu Ile Ser Glu Glu Glu Ile Pro Phe Lys Asp Asp Pro Arg Asp. 1 5

Glu Thr Tyr Lys Pro His Leu Glu Arg Glu Thr Pro Lys Pro Arg Arg 20

Lys Ser Gly Lys Val Lys Glu Glu Lys Glu Lys Lys Glu Ile Lys Val 35 40

2

Glu Val Glu Val Glu Val Lys Glu Glu Glu Msn Glu Ile Arg Glu Asp 50 50 Glu Glu Pro Pro Arg Lys Arg Gly Arg Arg Arg Lys Asp Lys Ser 75 80

15

Pro Arg Leu Pro Lys Arg Arg Lys Lys Pro Pro Ile Gln Tyr Val Arg 85  $\,$  90  $\,$  95

20

Cys Glu Met Glu Gly Cys Gly Thr Val Leu Ala His Pro Arg Tyr Leu 100

Gln His His Ile Lys Tyr Gln His Leu Leu Lys Lys Lys Tyr Val Cys 115

25

Pro His Pro Ser Cys Gly Arg Leu Phe Arg Leu Gln Lys Gln Leu Leu 130 8

Arg His Ala Lys His His Thr Asp Gln Arg Asp Tyr Ile Cys Glu Tyr 145 160

Ser His Asn Leu Ala Val His Arg Met 170 175 Cys Ala Arg Ala Phe Lys Ser 35

Ile His Thr Gly Glu Lys His Tyr Asn Val Arg Ser Val Asp Leu Leu 180

Val Asp Lys Arg His Leu Leu Ile Gly Thr Xea 195

8

(2) INFORMATION FOR SEQ ID NO: 377:

45

(A) LENGTH: 29 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 377:

(xi

Leu Ala Ser Phe Trp Gly Phe Thr Leu Arg Ala Ser Phe 20

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SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 136 amino acids

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INFORMATION FOR SEQ ID NO: 378:

WO 98/39448

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ઝ 45 6 35 25 20 5 5 50 Met Phe Asp Ser Leu Ser Tyr Phe Lys Gly Ser Ser Leu Leu Leu Met 1 5 10 Phe Tyr Asn His Lys Phe Leu Xaa 130 · 135 잗 Met Lys Lys Gly Lys Glu Leu Phe Ile Gln Gln Glu Arg Phe Phe Leu Asn 85 90 95 Asp Ser Phe Asn Glu Val 50 Tyr Leu His Asn His Ser Tyr Ala Ser Ile Gln Ser Asp Asp Leu Trp 35Leu Lys Thr Tyr Leu Ser Glu Asp Val Phe Gln His Ala Val Val Leu  $20 \ 30$ Gly Leu Met Ala Trp Arg Arg Glu Pro Ala Ser Gly Leu Ala Ala Cys Trp
1 10 15 (2) INFORMATION FOR SEQ ID NO: 379: Met Lys Thr Trp Thr Leu Gln Lys Gly Phe Pro Leu Val Thr Val Gln 65 70 70 75 Ser Cys His Leu 115 Lys Pro Glu Ile Gln Pro Ser Asp Thr Arg Tyr Met Pro Ser Phe  $100\,$ Ξ ξ. Ser Ser Gly Ser Arg Pro Trp 35 40 SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: 379: (A) LENGTH: 41 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Phe Cys Thr Leu Arg Trp Lys Tyr Phe Glu Val 120 125 Thr Asn Gln Thr Leu Asp Val Lys Arg Met 55

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Val Thr Lys Lys Ile Lys Val Tyr Glu Tyr Asp Thr Val Ile Gln Asp 180

Ser Ser Ile Glu Phe Asp Arg Asp Cys Asp Tyr Phe Ala Ile Ala Gly 175

Ala Val

Asp Ile His Tyr Pro Glu Asn Glu Met Thr Cys Asn Ser Lys 195 205

45

Ile Ser Cys Ile Ser Trp Ser Ser Tyr His Lys Asn Leu Leu Ala Ser 210 220

Ser Asp Tyr Glu Gly Thr Val Ile Leu Trp Asp Gly Phe Thr Gly Gln 235 230

Arg Ser Lys Val Tyr Gln Glu His Glu Lys Arg Cys Trp Ser Val Asp 255 255

Phe Asn Leu Met Asp Pro Lys Leu Leu Ala Ser Gly Ser Asp Asp Ala 260 265

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Lys Val Lys I 275

Leu Trp

Ser Thr

Asn Leu Asp Asn Ser Val Ala Ser Ile 280 285

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Glu Ala Lys Ala Asn Val Cys Cys Val Lys Phe Ser Pro Ser Ser Arg

8 5 25 7 30 Glu Glu Met Ser Gly Leu Tyr Ser Pro Val Ser Glu Asp Ser Thr Val 35 Oln Ile Oln Lys Glu Leu Ser Val Leu Glu Glu Asp Ile Lys Arg Val 20 25 Met Glu Phe Leu Lys Val Ala Arg Arg Asn Lys Arg Glu Gln Leu Glu 1 15 Pro Gln Phe Glu Ala Pro Ser Pro Ser His Ser Ser Ile Ile Asp Ser 50 55 Lys Gln Pro Trp Tyr Asn Ser Thr Leu Ala Ser Arg Arg Lys Arg Leu 90 95 Gln Glu Cys Leu Ser Lys 130 Ser Arg Ile Ser Asp Asp Ser Arg Thr Ala Ser Gln Leu Asp Glu Phe 115 120 125 The Ala His Phe Glu Asp Leu Glu Gln Cys Tyr Phe Ser The Arg Met 100 105 The Glu Tyr Ser Gln Pro Pro Gly Phe Ser Gly Ser Ser Gln The  $65\,$ Ala Thr Leu Ser Tyr Ala Ser Asp Leu Tyr Asn Gly Ser Ser Ile Val 145 150 (xi) SEQUENCE DESCRIPTION: (D) TOPOLOGY: linear (A) LENGTH: 468 amino acids
(B) TYPE: amino acid Phe Thr Arg Tyr Asn Ser Val Arg Pro Leu 135 SEQ ID NO: 380: 80 80

SEQUENCE DESCRIPTION: SEQ ID NO: 378:

(1) SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 380:

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55 2 39 8 45 S 15 2 25 35 Met Arg Lys Glu Asp Gly Phe Trp Phe Phe Phe Leu Phe Phe he le Leu Arg Ser Phe Lys Gly His Ile Asn Glu Lys Asn Phe Val Gly Leu 370 Lys Pro Tyr Cys 365 Ala Ser Asn Gly Asp Tyr Ile Ala Cys Gly Ser Glu Asn Asn Ser Leu 185 Tyr Leu Tyr Tyr Lys Gly Leu Ser Lys Thr Leu Leu Thr Phe Lys Phe 410 Phe Val Ser Ala Val Cys Trp Arg Ala Leu Pro Asp Gly Glu 415 Asn Vai Leu Ile Ala Asn Ser Gln Gly Thr Ile Lys Val Leu 450 Tyr His Leu Ala Phe Gly Cys Ala Asp His Cys Val His Tyr Tyr Asp 310 315 Lys Gln Pro Ile Met Val Phe Lys Gly His Arg Lys 325 335 Ala Val Ser Tyr Ala Lys Phe Val Ser Gly Glu Glu Ile Val Ser Ala 340 Lys Ser Val Leu Asp Lys Asp Arg Lys Glu Asp Asp Thr 420 Val Val Gly Ser Lys Phe Val Asn Gly Asn Lys Leu Val 20 (A) LENGTH: 29 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 382: SEQUENCE DESCRIPTION: SEQ ID NO: 381: Asp Ser Gln Leu Lys Leu Trp Asn Val Gly 355 8 (A) LENGTH: 29 amino acids (1) SEQUENCE CHARACTERISTICS SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 382: (2) INFORMATION FOR SEQ ID NO: 381: TYPE: amino acid 295 Leu Arg Asn Thr Leu Val Xaa ž Asp Thr Val Ξ Asn Glu Ser Thr 290 Glu 465 Ser 55 S <del>\$</del> 45 2 20 23 8 35 2

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582

Met Pro Leu Ala Pro Tyr Cys Asp Leu Leu Val Ala Leu Ser Phe Ala 1 5 15 15

Ser Ser Asp Phe Thr Leu Val Leu Glu Ser Pro Val Asp

(2) INFORMATION FOR SEQ ID NO: 383:

(A) LENGTH: 138 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 383: (D) TOPOLOGY: linear

Met Asn Ser Leu Val Ser Trp Gln Leu Leu Leu Phe Leu Cys Ala Thr  $_{\rm 1}$ 

His Phe Gly Glu Pro Leu Glu Lys Val Ala Ser Val Gly Asn Ser Arg \$20\$

Pro Thr Gly Gln Gln Leu Glu Ser Leu Gly Leu Leu Ala Pro Gly Glu 35 Thr Glu Arg Lys Pro Ala Ala Thr Ala Arg Leu 55 Gln Ser Leu Pro Cys 50

Ser Arg Arg Gly Thr Ser Leu Ser Pro Pro Pro Glu Ser Ser Gly Ser 65 75 80

Pro Gin Gin Pro Gly Leu Ser Ala Pro His Ser Arg Gin Ile Pro Ala 90 95

Pro Gln Gly Ala Val Leu Val Gln Arg Glu Lys Asp Leu Pro Asn Tyr 110

Asn Trp Asn Ser Phe Gly Leu Axg Phe Gly Lys Axg Glu Ala Ala Pro 125 Gly Asn His Gly Arg Ser Ala Gly Arg Gly 130

INFORMATION FOR SEQ ID NO: 384: 3

(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 384: (A) LENGTH: 74 amino acids

Met Ser Cys Phe Ile Asp Ser Xaa Asp Ser Lys Ile Leu His Leu Leu 1 5 10 15 15

Val Val Ser Phe lle Cys Xaa Leu Phe Leu Leu Ile Leu Thr His Gly 25

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583

Met Ser Ala Gly Glu Val Glu Arg Leu Val Ser Glu Leu Ser Gly Gly 1 15 Arg Glu Asp Phe Phe Val Leu Pro Xaa Ala 65 ( 70 Agn Ile Leu Ile Leu Arg Xaa Phe Phe Ser Val Xaa Xaa His Ser Leu Lys 35 40 Ile Glu Asp Glu Thr Ala Glu Asn Gly Val Pro Lys Pro Lys Val Thr 50 55 (2) INFORMATION FOR SEQ ID NO: 385: Val His Val Thr Ile Gly Asp Ile Lys Thr Gly Ala Pro Gln Tyr Gly 95 90 95 Glu Thr Glu Asp Asp Ser Asp Ser Asp Ser 65 Val Glu Arg Pro Glu Glu Glu Asn Ala Ser Ala Asn Pro Pro Ser Gly
35
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45 Phe Asn Tyr Gly Phe Asn Glu Asp Thr 165 Pro Gly Ser Ile Asn Gly Val Pro Leu Leu Glu Val Asp Leu Asp Ser 130 140 Val Tyr Gly Thr Thr Gly Thr Lys Val Lys Gly Val Asp Leu Asp Ala 115 Ser The Gly Gly Phe Glu Asp Lys Pro Trp Arg Lys Pro Gly Ala Asp Leu Ser Asp Tyr 145 150 Glu Thr Ala LeuiPro Ser Thr Lys Ala Glu Phe Thr Ser Pro Pro Ser 210 215 Thr Asn Lys Ile Thr Val Gln Gln Gly Arg Thr Gly Asn Ser Glu Lys 195 200 Gln Lys Asn Leu Glu Glu Tyr Leu Ile Leu Met Asn Lys Ala Leu Leu Thr 50 55 Tyz Gly Thr Ala Pro Val Asn Leu Asn Ile Lys Thr Gly Gly Arg  $100\,$ Ξ <u>X</u> Arg Ile Arg Met Gly Leu Glu Val Ile Pro Val Thr Ser Thr 180 185 ) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 521 amino a.

(B) TYPE: amino acid

(D) TOPOLOGY: linear Asp Glu Glu Glu Glu Trp Leu Tyr Gly Asp Glu Asn Glu 20 25 SEQUENCE DESCRIPTION: SEQ ID NO: 385: Trp Lys Ala Tyr Cys Glu Lys 170 175 Asp Asp Asp Glu Asp Asp 75

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5 25 20 2 30 8 35 55 8 3 Thr Glu Val Asp Asn Asn Phe Ser Lys Pro Pro Pro Phe Phe Pro Pro 275 280 285 Arg Ala As<br/>n Glu Asn Ser Asn Ile Gl<br/>n Val Leu Ser Glu Arg Ser Ala 260 265 270 Asp Val Ile Gly Gln Thr Ile Thr Ile Ser Arg Val Glu Gly Arg Arg 245 250 255 Leu Phe Lys Thr Gly Leu Pro Pro Ser Arg Arg Leu Pro Gly Ala Ile 225 230 230 235 Gly Ala Pro Pro Thr His Leu Pro Pro Pro Pro Phe Leu Pro Pro Pro 290 295 300 Gly Asn Val Ala Phe Pro His Leu Pro Gly Ser Ala Pro Ser Trp Pro 365 Gly His Ser Ser Gly Tyr Asp Ser Arg Ser Ala Arg Ala Phe Pro Tyr 340 345 Pro Pro Thr Val Ser Thr Ala Pro Pro Leu Ile Pro Pro Pro Gly Phe Pro 305 310 315 Glu Lys Glu Glu Arg His Arg Glu Glu Glu 450 450 460 Tyr Arg Glu Tyr Ala Glu Arg Gly Tyr Glu Arg His Arg Ala Ser Arg 435 His Ser Pro Thr Pro Ser Val Phe 420 Arg Asp Arg Glu Arg Glu Arg Thr Arg Glu Arg Glu Arg Glu Arg Asp 415 Lys Asp Arg Asp Arg Glu Arg Asp Arg Asp Arg Glu Arg Asp Arg Asp 385 390 Ser Glu Glu Gly Asp Ser His Arg Arg His Lys His Lys Lys Ser Lys Arg 495 Thr Arg His Lys Ser Ser Arg Ser Asn Ser Arg Arg Arg His Glu Ser 475 470 Ser The Glu Ala The Pro Ala Glu Xaa 515 520 Ser Lys Glu Gly Lys Glu Ala Gly Ser Glu 500 505 Leu Val Asp Thr Ser Lys Gln Trp Asp Tyr Tyr Ala Arg Arg Glu 370 380 Pro Pro Gly Ala Pro Pro Pro Ser Leu Ile Pro Thr Ile Glu Ser 330 335 Asn Ser Asp Glu Glu Arg Tyr Arg 425 430 Pro Ala Pro Glu Gln Glu 510

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INFORMATION FOR SEQ ID NO: 386: 3

SEQUENCE CHARACTERISTICS 3

(A) LENGTH: 137 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 386: ž Met Asn Ser Arg Gly lle Trp Leu Ala Tyr lle lle Leu Val Gly Leu 2

Leu His Met Val Leu Leu Ser Ile Pro Phe Phe Ser Ile Pro Val Val 20

Trp Thr Leu Thr Asn Val Ile His Asn Leu Ala Thr Tyr Val Phe Leu 15 2

His Thr Val Lys Gly Thr Pro Phe Glu Thr Pro Asp Gln Gly Lys Ala 50 2

Arg Leu Leu Thr His Trp Glu Gln Met Asp fyr Gly Leu Gln Phe Thr 65

ŝ 3 S Ser Ser Arg Lys Phe Leu Ser Ile Ser Pro Ile Val Leu Tyr 90 85

22

Ala Ser Phe Tyr Thr Lys Tyr Asp Ala Ala His Phe Leu Ile Asn Thr 100

Ala Ser Leu Leu Ser Val Leu Leu Pro Lys Leu Pro Gln Phe His Gly 115 9

Val Arg Val Phe Gly Ile Agn Lys Tyr 130

35

(2) INFORMATION FOR SEQ ID NO: 387:

(A) LENGTH: 186 amino acids (B) TYPE: amino acid SEQUENCE CHARACTERISTICS: £

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SEQUENCE DESCRIPTION: SEQ ID NO: 387:

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Met Ala Ala Gln Lys Asp Gln Gln Lys Asp Ala Glu Ala Glu Gly Leu 1 5 15 45

Thr Thr Leu Leu Pro Lys Leu Ile Pro Ser Gly Ala Gly Arg 20 Glu Trp Leu Glu Arg Arg Arg Ala Thr Ile Arg Pro Trp Ser Thr Phe 45 46 Ser Gly 20

Asp Gin Gin Arg Phe Ser Arg Pro Arg Asn Leu Gly Giu Leu Cys 50 60 Val 55

Gin Arg Leu Val Arg Asn Val Glu Tyr Tyr Gin Ser Asn Tyr Val Phe 65

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Val Phe Leu Gly Leu Ile Leu Tyr Cys Val Val Thr Ser Pro Met Leu 85  $\,$  95  $\,$ 

586

Leu Ala Val Phe Ghy Ala Cys Tyr Ile Leu Tyr Leu 100 110 Leu Val Ala

Arg Glu Val Ser Pro 125 Leu Glu Ser Lys Leu Val Leu Phe Gly 115 Arg Thr

Phe Pro Phe Phe Trp 140 Leu Ala Gly Ala Gly Ser Ala Val Phe Trp Val Leu Gly Ala Thr Leu 145 Ala His Gln Tyr Ala Leu Ala Gly Gly Ile Ser 130

2

Val Val Ile Gly Ser His Ala Ala Phe His Gln Ile Glu Ala Val Asp 175

2

Gly Glu Glu Leu Gln Met Glu Pro Val Xaa 180 185 2

(2) INFORMATION FOR SEQ ID NO: 388:

(i) SEQUENCE CHARACTERISTICS:

23

(A) LENGTH: 1 emino acids TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 388:

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(2) INFORMATION FOR SEQ ID NO: 389:

(i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 389: (B) TYPE: amino acid (D) TOPOLOGY: linear

Mat Leu Ser lle Phe Tyr Phe Ala Ile Pro Val Gly Ser Gly Leu Gly 1 5 15 15 45

Tyr ile Ala Gly Ser Lys Val Lys Asp Met Ala Gly Asp Trp His Trp 20

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Ala Leu Arg Val Thr Pro Gly Leu Gly Val Val Ala Val Leu Leu Leu 45 Leu Val Val Arg Glu Pro Pro Arg Gly Ala Val Glu Arg His Ser 50 60 Phe

Trp Ala Asp Leu Arg Ala Asn Pro Thr Ser Trp 70 Ž Pro Pro A 20

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Leu Ala Arg Asn Pro Ser Phe Val Leu Ser Ser Leu Gly Phe Thr Ala 8

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Val Ala Phe Val Thr Gly Ser Leu Ala Leu Trp Ala Pro Ala Phe Leu 100 105 85

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λg Ser Arg Val Val Leu Gly Glu Thr Pro Pro Cys Leu Pro Gly 115 120

Leu Thr Gly Val Leu Gly Val Gly Leu Gly Val Glu Ile Ser Arg Arg 145 150 150 Ser Cys Ser Ser Ser Asp Ser Leu Ile Phe Gly Leu Ile Thr Cys 130 140

5

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Leu Arg His Ser Asn Pro Arg Ala Asp Pro Leu Val Cys Ala Thr Gly
175
175

5

Gly Ser Leu Leu Gly Ser Ala Pro Phe Leu Phe Leu Ser Leu Ala Cys Ala Arg 180 185 Ile Val Ala Thr 195 Tyr Ile Phe Ile Phe Ile Gly Glu Thr Leu 200

20

Phe

Val Leu Leu Phe Val Cys Phe Cys Gly Leu Ser Tyr Val Val Ile 35  $40\,$ 

25

Ala Gln Val Pro Val Gly Leu Leu Cys Ile Thr Glu Xaa 50 55 60

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Met Tyr Val Asn Tyr Gly Thr Arg Asn Tyr Ser Thr Glu Gly Pro Ala 1 5

Ala Leu Leu

Asp Gln Ala Lys Leu Ser Leu Leu Val Trp Val Leu Cys 20 25 30

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SEQUENCE DESCRIPTION: SEQ ID NO: 391:

(2) INFORMATION FOR SEQ ID NO: 391:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 61 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

20

25 Leu Ser Met Asn Trp Ala Ile Val Ala Asp Ile Leu Tyr Val Val 210 215 220

Ile Pro Thr . 225 ₽rg Arg Ser Thr Ala Glu Ala Phe Gln Ile Val Leu Ser 230 235

His Leu Leu Gly Asp Ala Gly Ser Pro Tyr Leu Ile Gly Leu Ile Ser 245 250 255

30

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SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(2) INFORMATION FOR SEQ ID NO: 392:

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Val Ala Phe Val His Ile Ser His Ile Gly Val Arg Gln Val Asn Thr . 20 25

Met Leu Trp Phe Ala Asn Phe Phe Thr Tyr Leu Phe Leu Ser Gin Ser 1 10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 392:

3

Lys Lys Lys Ile Pro Ala Gly Arg Xaa Leu Phe Pro Phe Gly 65 70

Asn Cys Tyr Phe Ser Arg Lys Ser Tyr Cys Tyr Gly Ile Leu Asn Pro 35

50

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SEQUENCE CHARACTERISTICS:

INFORMATION FOR SEQ ID NO: 393:

30

₽ Leu Arg Arg Asn Trp Pro 260 Pro Ser Phe Leu Ser Glu Phe Arg 265 270

Ala Leu Gln Phe Ser Leu Met Leu Cys Ala Phe Val Gly Ala Leu Gly 275  $280\,$ 

35

8 Gly Ala Leu Pro Gly His Arg His Leu His Xaa 290 295

(2) INFORMATION FOR SEQ ID NO: 390:

25

Ξ SEQUENCE CHARACTERISTICS:

(A) LENGTH: 49 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Gly Pro Gln Gly Trp Val Arg Pro Leu Lys Thr Ala Pro Lys Leu 1 , 5 Ě SEQUENCE DESCRIPTION: SEQ ID NO: 390:

50

55 Gly Glu Ala Ile Arg Leu Ile Leu Phe Leu Asn Phe Val Lys Gln Cys 20 25

Ile Ala Ser Val Asn Leu Cys Ile Leu Arg Leu Asn Ile Thr Pro Leu 35 1 40

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 393:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 36 amino acids

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289

Met Pro Gly Ala Phe Ser Glu Thr Val Ile Asn Asp Leu Leu Ser Leu  $_{\rm 1}$ 

Ser Tyr Ser Thr Leu Ser Gly Val 25 Pro Ala Glu Leu 3 % Phe Leu Val

Tyr Arg Asn Ala 35

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(2) INFORMATION FOR SEQ ID NO: 394:

SEQUENCE CHARACTERISTICS: 3

2

(A) LENGTH: 180 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 394:

Met Ala Gin Ser Arg Asp Gly Gly Asn Pro Phe Ala Glu Pro Ser Glu 8

Leu Asp Asn Pro Phe Gln Asp Pro Ala Val Ile Gln His Arg Pro Ser

Arg Gln Tyr Ala Thr Leu Asp Val Tyr Asn Pro Phe Glu Thr Arg Glu 35

22

Leu Pro Pro Pro 9 S Pro Pro Pro Ala Tyr Glu Pro Pro Ala Pro Ala ဂ္ဂ ജ Gln Pro Ser Arg Lys Leu Ser Pro Thr Glu Pro 70 75 80 Ser Ala Pro Ser 8 Lys Asn Tyr Cly Ser Tyr Ser Thr Cln Ala Ser Ala Ala Ala Thr 95 35

Ala Glu Leu Leu Lys Lys Gln Glu Glu Leu Asn Arg Lys Ala Glu Glu 100

Arg Arg Ser Glu Ser Cys Ser Met Leu Pro Trp Xaa Ala Gln 115 Leu Asp 6

Leu Asp Arg Thr lle Gly Pro Leu Tyr Leu Leu Phe Val Gln Phe 130 3 5

Pro Ala Phe Ser Arg Thr Ser Pro Trp Arg Ser Pro Lys Asn Phe 150 Ser 145 Arg Arg Leu Tyr Pro Pro Cys Thr Thr Ser Cly Cys Ala Ala Arg Trp 175 Phe Ser Xaa 180 Xaa

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(2) INFORMATION FOR SEQ ID NO: 395:

(1) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 21 amino acids

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 395: (D) TOPOLOGY: linear

Met Pro Thr Pro Cys Thr Ser Leu Pro Ser Cys Cys Gln His Arg Ser

lle Thr Met Thr Leu

2

(2) INFORMATION FOR SEQ ID NO: 396:

15

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 60 amino acids
(B) TYPE: amino acid 3

(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 396: Ī

20

Pro Leu Phe Ile Pro Leu Ile Phe Phe Leu Ser Leu Leu His Cys 10 Met

Lys His Pro Ile Gin Met Ser Leu Cys Met Cys Val Asn Ile 20 Ser g

23

Leu Val Trp Ser Pro Val Arg Trp lle Phe Cly Ser Lys Cly Leu 35 45 Ser

Val His Leu Gln Ser Ser Gln Arg Pro Ser 55 60 Phe

8

(2) INFORMATION FOR SEQ ID NO: 397:

35

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 152 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 397:

Met Ala Gly Pro Arg Pro Xaa Trp Arg Aap Gln Leu Leu Phe Met Ser  $_{1}$  10  $_{5}$ 45

ile ile Val Leu Val ile Val Val ile Cys Leu Met Leu Tyr Ala Leu 20 30

Leu Trp Glu Ala Gly Asn Leu Thr Asp Leu Pro Asn Leu Arg Ile Gly 35 င္တ

Phe Tyr Asn Phe Cys Leu Trp Asn Glu Asp Thr Ser Thr Leu Gln Cys 50 60

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His Gln Phe Pro Glu Leu Glu Ala Leu Gly Val Pro Arg Val Gly Leu 65 79 Gly Leu Ala Arg Leu Gly Val Tyr Gly Ser Leu Val Leu Thr Leu Phe \$95\$

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591

20 5 4 25 5 50 5 35 30 S 55 Pro Ser Arg Gly Leu 145 Gly Gly Leu Gly Leu Phe Leu Ser Tyr Val Trp Asn Gly Ser Xaa Ser 130 140 Trp Arg Leu Ala Val Gly Phe Leu Ala Val Ser Ser Val Leu Leu Ala 115 120 125 Ala Pro Gln Pro Leu Leu Leu Ala Gln Cys Asn Xaa Asp Glu Arg Ala 100 105 · Xaa Gly Xaa Xaa Pro Ala Glu Arg Xaa Arg His Gln Pro Pro Gln Pro 35 40 45 Met Ser Asp Gly Phe Asp Arg Ala Pro Gly Ala Gly Arg Gly Arg Xaa 1 10 15 Thr Pro Pro Pro Gly Ala Gln Cys Glu Val Pro Ala Ser Pro Gln Arg 65 70 75 Lys Ala Pro Gly Phe Leu Gln Pro Xea Pro Leu Arg Gln Pro Arg Thr 50 55 Arg Gly Leu Gly Arg Gly Gly Gly Gly Bro Xaa Gly Gly Gly Phe Pro 20 25 30 Met Ala Lys Pro Gin Val Val Val Ala Pro Val Leu Met Sex Lys Leu 115 120 125 Pro Ser Arg Pro Gly Ala Leu Pro Glu Gln Thr Arg Pro Leu Arg Ala 95 5 Glu Thr Glu Ile Glu Gln Phe Ala Glu Thr Leu Asn Gly Cys Val Thr 180 185 The Glu Ser Tyr Glu Asp Gly Cys Glu Asp Tyr Pro The Leu Ser Glu 145 150 Ser Val Asn Ala Pro Glu Phe Tyr Pro Ser Gly Tyr Ser Ser Ser Tyr 130 140 Pro Tyr Val Gln Asp INFORMATION FOR SEQ ID NO: 398: £ ž Ser SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 398: Ser Gln Asp Lys Ile Pro Gln Gln Asn Ser Glu Ser Ala 100 105 110 (A) LENGTH: 480 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Phe Leu Asn His Leu Thr Glu Gln Pro Gly Ser Phe 165 170 175 1 Gly Phe Xaa 150

> 23 20 2 5 8 35 30 5 50 Asn Ala Leu i 305 Leu Leu Gin Arg Cys Arg Thr Glu Tyr Glu Val Lys Asp Gin Ala 255 245 Leu Ser His His Leu ' 붛 Gln Val Thr Arg Ala Asp Ile Leu Gln Val Gly Leu Arg Glu Leu Leu 290 300 Phe Leu Gly Glu Leu Tyr Leu Asn Leu Glu Ile Lys Gly Thr Asn Gly 275 280 ala Lys Gly Asp Glu Val Thr Arg Lys Arg Phe His Ala Phe Val Leu 265 265Val Leu Asp Ala Asn Cys Ser Arg Asp Val Lys Gln Met Leu Leu Lys 355 360 Lys Gly Lys Met Asp Met Glu Glu Ile Ile Gln Axg Ile Glu Asn Val  $340^\circ$ Lys Leu Leu Lys Leu Thr Gly Ser Val Leu Glu Asp Ala Trp Lys Glu 325 330 335 Glu Pro Thr Phe Tyr Thr Ser Asp Gly Val Pro Phe Thr Ala Ala Asp 415 Leu Val Glu Leu Arg Ser Ser Asn Trp Gly Arg Val His Ala Thr Ser 370 375 380 Phe Thr Tyr Arg Glu Ala Thr Pro Glu Asn Asp Pro Asn Tyr Phe Met 385 Ala Tyr Glu Lys Phe Cys Leu Glu Ser Glu Arg Lys Arg Lys Gln Xaa 465 470 475 Pro Pro Ser Ile Pro Asn Phe Ser Tyr Met Gly Ala Arg Leu Cys Asn Tyr 210 215 220 253 Pro Asp Tyr Glu Glu Asn Gly Thr Asp Leu Ser Gly Ala Gly Asp 435 Asp Tyr 195 Leu Asp Asp Ile Asp Asp Glu Met Asp Pro Glu Ile Glu Glu 455 Phe Ser Asn Pro Met Asp Asp Asn Leu Ile Cys Ala Val 310 320 Gln Glu Lys Tyr Gln Glu Leu Clu Arg Glu Asp Pho 420 425 Thr Ile Ser Pro Gln Ser Gly Asn Phe Arg Gln 230 235 240 200 205 400 400

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INFORMATION FOR SEQ ID NO: 399:

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Thr Asp Asp Ala Leu Gln Glu Leu Val Glu Leu Ile Tyr Gln Gln Ala

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Trp Gln Asp Trp Ala Leu Gly Val Leu His Ala Lys Ile Ile Ala Ala 275 Ala Gly Val Val Pro Leu Leu Leu Gly Leu Leu Phe Glu Leu Val Ile 255 Val Ala Pro Leu Arg Val Pro Leu Asp Gln Thr Pro Leu Phe Tyr Pro 260. Glu frp Ser Leu Met Ile Met Lys Thr Leu Ile Val Ala Val Leu Leu 230 235 Leu His 80 Ser Leu Ile Cys Leu Thr Leu Pro Val Phe Ala Gly Arg Trp Leu Met 175 175 Leu Tyr Val Cys Trp Leu Thx lle Axg Ala Val Thx Val Met 195 Val Ala Trp Met Pro Gln Gly Arg Arg Val 11e Phe Gln Lys Val Lys 210 Arg Ile Phe Leu Leu Ile Val Phe Met Cys Ile Thr Leu Leu Ile Ala 145 Ser Fhe Try Thr Gly Thr Ala Lys Ile Hie Glu Leu Tyr Thr Ala Ala 185 Met Glu Pro Lys Thr Ile Thr Asp Ala Leu Ala Ser Ser Ile Ile Lys Leu Pro Asn Phe Leu Pro Tyr Asn Val Met Leu Tyr Ser Asp 20 Ala Pro Val Ser Glu Leu Ser Leu Glu Leu Leu Leu Leu Gln Val Val 35 Pro Ala Leu Leu Glu Glu Gly His Thr Arg Gln Trp Leu Lys Gly 50 60 Gin Val Asn Asn Gin His Ala Arg Asn Asn Asn Ala Ile Pro Val 105 Val Gly Glu Gly Leu His Ala Ala His Gln Ala Ile Leu Gln Gln Gly 115 Gly Pro Val Gly Phe Gln Xaa Tyr Arg Arg Pro Leu Asn Phe Pro Leu 130 Leu Val Arg Ala Trp Thr Val Thr Ala Gly Tyr Leu Leu Asp. 75 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 399: (A) LENGTH: 423 amino acids SEQUENCE CHARACTERISTICS TYPE: amino acid (B) TYPE: amino acid (D) TOPOLOGY: linear 3 Cys Gly Ser Val Ę 55 တ္သ 35 <del>6</del> 5 2 25 ഉ 2 2

Leu Val Asn Tyr Glu Arg Lys Ser Gly Lys Gln Gly Ser Ser Pro Pro 405 415 Vel Tyr Ala Asn Gly Ile Arg Asn Ile Asp Leu His Tyr Ile Val Arg 310 315 Lys Leu Ala Ala Pro Val Ile Ser Val Leu Leu Leu Ser Leu Cys Val 336 335 Leu Met val val Leu Met Ala Ile Leu Ser Phe Gln val Arg Gln Phe Lys 370 Arg Leu Tyr Glu His Ile Lys Asn Asp Lys Tyr Leu Val Gly Gln Arg 185 Pro Tyr Val Ile Ala Ser Gly Val Val Pro Leu Leu Gly Val Thr Ala 345 lle Thr Leu Met Gly Pro Gln Trp Trp Leu Lys Thr Val Ile Glu Gln 290 190 Σeς Glu Met Gln Asn Leu Val His Arg Arg Ile Tyr Pro Phe 360 Pro Pro Gln Ser Ser Gln Glu 420 9 2 8 22

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 400: (A) LENGTH: 78 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 400: 62 8 32

Asp Gly Ala Leu Ile Tyr Arg Lys Leu Leu Phe Asn Pro Ser Gly Pro 50Leu Thr Val Gly Gly Gly Val Phe Ala Leu Val Thr Ala Val Cys Cys Leu Ala 45 45 Leu Arg Leu Asp Ile Ile Asn Ser Leu Val Thr Thr Val Phe Met 10Leu ile Val Ser Val Leu Ala Leu ile Pro Glu Thr Thr Thr 25 20 20 Met 45 <del></del>

Tyr Gln Lys Lys Pro Val His Glu Lys Lys Glu Val Leu Xaa 65

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(2) INFORMATION FOR SEQ ID NO: 401:

22

(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 74 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 401:

S His Cys Trp Gly Leu Pro Leu His Val Ala Pro Leu Cys Arg Gly His 20 25 Met Leu Lys Gln Val Met Phe Val Phe Ser Gly Met Gly Pro Arg Ser 1 10

5 Gln Ala Asp Ser Ser His Leu Leu Pro Leu Lys His Gln Gly Ala Trp 35

Asn Arg Asn Leu Ala Asn Gln Arg His Phe Phe Cys Pro Ser Ile Phe 50 55

His The Cys Pro 才 Val Leu Phe Phe Xaa 70

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E SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO:

402:

χĽ SEQUENCE DESCRIPTION: SEQ ID NO: 402:

Ala Arg Thr Ile Leu Val Leu Tyr Leu Ser Leu Gln Arg Leu Glu Asn 1 5

Leu Ala Tyr His 20

30

(2) INFORMATION FOR SEQ ID NO: 403:

35

Ξ SEQUENCE CHARACTERISTICS:

χĹ SEQUENCE DESCRIPTION: SEQ ID NO: 403: (A) LENGTH: 87 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

6

Met Pro Leu Pro Ser Val Pro Ile Leu Gly Ile Phe Ser Phe Leu Ile 1 5

3

Pro. Ser Ser Gln Gly Val Ser Tyr Thr Lys Leu Pro Ile Ser Ser Pro 20 25

8 Gln Tyr Ser Pro Phe Val Asn Asp His Phe Ser Phe Leu Asn Pro Phe 35  $40\,$ 

55 Pro Val Gln Ile His Thr Gly Phe Ala Arg Val Gly Ser Tyr Met Gln 50 55

Met Pro Leu Val His Leu Cys Leu Leu Gln Thr Ser Leu Met Lys Asn 65 70 75

8 ser Gly Val Gln Gln Gly Ser 85

(2) INFORMATION FOR SEQ ID NO: 404: (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 92 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 404: (D) TOPOLOGY: linear (B) TYPE: amino acid

5 Met Asn Ala Ala Met Val His Ile Asn Arg Ala Leu Lys Leu Ile Ile 1 10 15

15  $\mbox{Arg Leù Phe Leu Val Glu Asp Leu Val Asp Ser Leu Lys Leu Ala Val <math display="inline">\mbox{20}$ 

Phe Met Trp Leu Met Thr Tyr Val Gly Ala Val Phe Asn Gly Ile Thr 35

Leu Leu Ile Leu Ala Glu Leu Leu Ile Phe Ser Val Pro Ile Val Tyr 50 60

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Gln Thr Lys Ser Ile Val Glu Lys Ile Pro Ser Lys 85 Glu Lys Tyr Lys Thr Gln Ile Asp His Tyr Val Gly Ile Ala Arg Asp 65 70 75

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3 INFORMATION FOR SEQ ID NO: 405:

(A) LENCTH: 21 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 405: (i) SEQUENCE CHARACTERISTICS:

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8 Met Ala Cys Ser Cys Leu Met Ile Gln Ser Phe Ser Thr Ser Ala Leu 1 5

Val Leu Phe Tyr Gly 20

45

(2) INFORMATION FOR SEQ ID NO: 406:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 174 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 406: (B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Glu Glu Gly Gly Asn Leu Gly Gly Leu Ile Lys Met Val His Leu 1 15

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Leu Val Leu Ser Gly Ala Trp Gly Met Gln Met Trp Val Thr Phe Val 20 25 10

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Ser Gly Phe Pro Ala Phe Pro Lys Pro Ser Pro Thr Tyr Leu Arg Thr 45

Ser Ala Glu Gln Thr Leu Pro Leu Leu Fro His Leu His Gly Leu 50 60

Leu His Gln Pro Leu His Leu Gly Phe Thr Ala Cys Leu Gly Ser  $$75\$ 8 3 2

Ala His Ile Leu Gly Gly Gln Pro Ala Leu Pro Ala Val Pro Glu Pro 95

Tyr Ala Gly His Cys Gln Arg Pro Leu Ala Gly Thr Pro His His Ser 100

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Gly Arg Tyr Gln Ala Ala Asn Arg Phe Pro Ile Leu Asn Ala Xaa Cys 130 Cys His Val Gly Pro Ala Asn Arg Gly Arg Arg Ser Glu Ala Trp Val 115

Ser Ala 160 Glu Arg Arg Thr Pro Ser Thr Val Leu Ser Ala Arg Ile Ser 145 23

Thr Met Gly Cys Pro Leu Phe Ala Ile Trp Ala Ala Ser Xaa 170

(2) INFORMATION FOR SEQ ID NO: 407:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 407: (D) TOPOLOGY: linear

Glu Gln Asn Ser Ala Thr Val Glu Pro Ser Ser His Glu Ile Leu His Met Ala Phe Ile Leu Leu Phe Tyr Cys Leu Met Thr Phe Leu Ser Leu 1 10 6

Leu Gln Asn Cys Phe Glu Leu Leu Arg Thr Ser Thr Ser Gln Cys 35 40 7 45

Glu Gly Ile Pro Cys Gln Arg Tyr Gln Asn Gly Leu His Ile Xaa 50 60 궕 S

(2) INFORMATION FOR SEQ ID NO: 408:

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(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 280 amino acids
(B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 408: (D) TOPOLOGY: linear

Met Glu Ala Val Val Asn Leu Tyr Gln Glu Val Met Lys His Ala Asp

Pro Arg Ile Gln Gly Tyr Pro Leu Met Gly Ser Pro Leu Leu Met Thr  $20 \ 25 \ 30$ 

Ser Ile Leu Leu Thr Tyr Val Tyr Phe Val Leu Ser Leu Gly Pro Arg 135 46 Ile Met Ala Asn Arg Lys Pro Phe Gln Leu Arg Gly Phe Met Ile Val $_{\rm 50}$ 

2

Tyr Asn Phe Ser Leu Val Ala Leu Ser Leu fyr Ile Val Tyr Glu Phe 65

2

Asp Tyr Ser Asn Ser Pro Glu Ala Leu Arg Met Val Arg Val Ala Trp 110 Leu Met Ser Gly Trp Leu Ser Thr Tyr Thr Trp Arg Cys Asp Pro Val. 85

2

Leu Phe Leu Phe Ser Lys Phe Ile Glu Leu Met Asp Thr Val Ile Phe. 115 is Leu Arg Lys Lys Asp Gly Gin Val Thr Phe Leu His Val Phe His 130

23

His Ser Val Leu Pro frp Ser Trp Trp Gly Val Lys Ile Ala Pro 145

8

Gly Gly Met Gly Ser Phe His Ala Met Ile Asn Ser Ser Val His Val. 175 35

Ile Met Tyr Leu Tyr Tyr Gly Leu Ser Ala Phe Gly Pro Val Ala Glni 180 Pro Tyr Leu Trp Trp Lys Lys His Met Thr Ala Ile Gln Leu Ile Gln 195, 195, 6

Phe Val Leu Val Ser Leu His Ile Ser Gln Tyr Tyr Phe Met Ser Ser 210 45

Lys Gly Lys Arg Leu Pro Arg Ala Leu Gln Gln Asn Gly Ala Pro Gly 260 Cys Asn Tyr Gln Tyr Pro Val Ile Ile His Leu Ile Trp Met Tyr Gly 235Thr ile Phe Net Leu Phe Ser Aan Phe Trp Tyr His Ser Tyr Thr 255 255 S

Ile Ala Lys Val Lys Ala Asn Xaa 280

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60 . (2) INFORMATION FOR SEQ ID NO: 409:

Ξ SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 284 amino acids

ž SEQUENCE DESCRIPTION: SEQ ID NO: 409:

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Met Xaa Leu Trp Pro Gln Thr Cys Ser Gly Lys Phe Asp Gly Thr Leu 5

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SEQUENCE DESCRIPTION: SEQ ID NO: 410:

(A) LENGTH: 187 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Leu Phe Leu Phe Phe Val Ile Ile Phe Leu 1 5 10

Phe Val Phe Leu 15

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 410:

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Ala Phe Ser Ile His Xaa Leu Ala Val Ile Leu Gly Asp 20 25 Gln Leu Thr 30

Ala Ala Asp Leu Val Pro Ile Phe Asn Gly Phe Leu Lys Asp Leu Asp 35  $40\,$ His Asp Phe Leu Lys Leu 60

15

Glu Val Arg Ile Gly Val Leu Lys His Leu 50 55

20 Leu His Ile Asp Lys Arg Arg Glu Tyr Leu Tyr Gln Leu Gln Glu Phe 65 70

25 Leu Val Thr Asp Asn Ser Arg Asn Trp 85 Arg Phe Arg Ala Glu Leu Ala 90 95

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Glu Leu Lys Gln Arg Phe Ser Val Phe Gly Glu Ile Glu Glu Cys Thr  $90\,$ 

arg Arg Val Val Phe Ile Gly Lys Ile Pro Gly Arg Met Thr Arg Ser 65 70 75 80

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His Tyr Gln Arg Gln Arg Val Leu Gln Lys Glu Arg Ala Ile Glu Glu 50 50

Gly Xaa Ser Asp Arg Arg Arg 35

Arg Tyr Ser Ser Tyr Arg Ser His Asp 40

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Leu Ile Ile Gln Phe Ser Lys Pro Leu Thx Asn Pro His Pro Pro Ala 20 25 30

Glu Gln Leu Ile Leu Leu Leu Glu Leu Tyr Ser Pro Arg Asp Val Tyr 100 105

Asp Tyr Leu Arg Pro Ile Ala Leu Asn Leu Cys Ala Asp Lys Val Ser 115 120

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Ser Val Arg Trp Ile Ser Tyr Lys Leu Val Ser Glu Met Val Lys Lys 130 135

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Leu His Ala Ala Thr Pro Pro Thr Phe Gly Val Asp Leu Ile Asm Glu 145

Leu Val Glu Asn Phe Gly Arg Cys Pro Lys Trp Ser Gly Arg Gln Ala 165 170

Phe Val Phe Val Cys Gln Thr Val Ile Glu Asp Asp Cys Leu 180 185 190 Pro Met

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Asp Gln Phe Ale Vel His Leu Met Pro His Leu Leu Thr Leu Ale Asn 195 200 205

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dsγ Arg Val Pro Asn Val Arg Val Leu Leu Ala Lys Thr Leu Arg Gln 210 220

50 Thr Leu Leu Glu Lys Asp Tyr Phe Leu Ala Ser Ala Ser Cys His Gln 240 225

Glu Ala Val Glu Gln Thr Ile Met Ala Leu Gln Met Asp 250 Arg Asp Ser 255

Asp Val Lys Tyr Phe Ala Ser Ile His Pro 260 265 Ala Ser Thr : Lys Ile Ser 270

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5 C Asp Ala Met \$er Thr Ala Ser Ser Thr Tyr Xaa 275 280

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XAA Cys Lys Arg Ser Tyr Ser Asp Leu Asp Ser Asn Arg Glu Asp Phe 145 150 150

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Tyr Ala Glu Glu Ala Phe Ala Ala Ile Glu Ser Gly His Lys Leu Arg 115 120 126

Gln Ala Asp Glu Gln Pro 130

Phe Asp Leu Cys 135

Phe Gly Gly Arg Arg Xaa 140

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Ile His

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Arg val Gln Gly Asp Asn Tyr Gly Phe Val Thr Tyr Arg 100 105

Asp Pro Ala Pro Val Lys Ser Lys Phe Asp Ser Leu Asp Phe Asp Thr 175

3 Leu Leu Lys Gln Ala Gln Lys Asn Leu Arg Arg 180

8 (2) INFORMATION FOR SEQ ID NO: 411:

E SEQUENCE CHARACTERISTICS acids

(A) LENGTH: 237 amino a (B) TYPE: amino acid (D) TOPOLOGY: linear

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Met Lys Leu Pro Gly Lys Phe Arg Arg Ala His Gln Gly Asn Leu Glu 1 15 SEQUENCE DESCRIPTION: SEQ ID NO: 411:

602

Ser Lys Ile Tyr 110

Phe

Ser

Thr 160

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Phe Glu Gln Leu Glu Tyr Trp Ala Asn Asn Phe Asp Asp Phe Ala Ala 65 Gly Asn Ser Ser Leu Ala Pro Ala Asn Gly Ser Ala Pro Cys Gly Ser 50 60 Ala Leu Val Thr Leu Trp Asn Leu Met Val Val Asn Asn Trp Gln Val 85 Leu Phe Arg Asp Ile Leu Glu Glu Pro Gly Glu Asp Glu 165 175 Leu Thr Glu Arg Leu Ser Gln His Pro His Leu Trp Leu Cys Arg Xoa 180 Asn Val Val Val Ala Phe Gly Leu Ile Leu Ile Ile Glu Ser Leu 1 10 11 15 15 15 16 16 17 Met Asn Trp Gly Leu Ser lle Trp Leu His Tyr Tyr Glu Lys Lys Lys 1 5 15 15 Phe Val Leu Trp Trp Leu Val Ser Ser Val Ile Trp Val Asn Leu 115 Leu Ala Leu Ile Leu Glu Asn Phe Leu His Lys Trp Asp Pro Arg 130 His Leu Gin Pro Leu Ala Gly Thr Pro Glu Ala Thr Tyr Gin Met 145 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 414: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 413: Phe Leu Asp Ala Tyr Arg Arg Tyr Ser Gly Pro Trp 100 (A) LENGTH: 51 amino acids (A) LENGTH: 21 amino acids (1) SEQUENCE CHARACTERISTICS: (i) SEQUENCE CHARACTERISTICS. (2) INFORMATION FOR SEQ ID NO: 414: (B) TYPE: amino acid INFORMATION FOR SEQ ID NO: 413: (B) TYPE: amino acid (D) TOPOLOGY: linear Gly Glu Gln Cys Pro Val Glu Leu 3 45 23 ଌ 2 15 2 ഉ 35 **4** 55 25 Ser Gln Leu Thr Ser Glu Ser Tyr Tyr Lys Glu Thr Leu Ser Val Pro  $26\ 26\ 30\ 30$ Lys Pro Met Ala Val Val Ala Ser Thr Val Leu Gly Leu Val Gln 15 Asn Met Arg Ala Phe Gly Gly Ile Leu Val Val Val Tyr Val Phe \$20\$Ala Ile Ile Gly Ile Asn Leu Phe Arg Gly Val Ile Val Ala Leu Pro Thr Val Glu His Ile Ile Gln Glu Leu Lys Asp Ile Phe Ser Glu Gln 35 Ser Asp Leu Pro Asn Pro Asp Thr Leu Ser Ala Glu Leu His Cys Trp 95 Ile Tyr Glu Ala Leu His Leu Pro Asp Ile Lys Phe Pro Asn Val 115 Tyr Ala Leu Leu Lys Val Leu Cys Ile Leu Pro Val Met Lys Val Glu 130 Asn Thr Leu Thr Asp Gln Arg Ser Ser Asn Leu Ala Leu Leu Asn Ile 175 His Leu Lys Ala Leu Lys Cys Leu Ser Leu Val Pro Ser Val Met Gly 50 60 Thr Ser Glu Glu His His Ala Asp Met Tyr Arg 70 75 80 Arg Ile Lys Trp Lys His Arg Gly Lys Asp Ile Glu Leu Pro Ser Thr 100 Asn Glu Arg Tyr Glu Asn Gly Arg Lys Arg Leu Lys Ala Tyr Leu Arg 145 185 Phe Asp Ile Lys His Asp Leu Asp Leu Met Val Asp Thr Tyr Ile 180 Leu Tyr Thr Xaa Xaa Ser Xaa Leu Xaa Thr Xaa Xaa Ser Xaa Xaa 195 Glu xaa xaa xaa xaa xaa xaa xaa xaa xaa Gly xaa xaa xaa xaa 210 Xaa Xaa Xaa Arg Glu Lys Ala Val Arg Cys Met Xaa 230 SEQUENCE DESCRIPTION: SEQ ID NO: 412: (A) LENGTH: 192 amino acids (i) SEQUENCE CHARACTERISTICS: INFORMATION FOR SEQ ID NO: 412: 215 Leu Lys Phe Asn X; Gln 65 Asn ş Val Asp 225 3 Met 45 8

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30 25 20 6 35 15 5 55 50 24 Ala Pro Xaa 50 Met Leu Ile Ile Ser Leu Arg Pro Gln Phe Pro Ser Leu Ile Val Gln 1 15 Glu Gln Val Phe Leu Val Ile Leu Ala His Val Val Arg Arg Cys Ala  $20 \ 30$ Phe His Pro Met Val Lys Val Cys Asn Asp Ser Asp Arg Trp Ser Leu Ile Ser Leu 1 15 Ser Asp Gly Ile Leu Gln Phe Glu Ser Ser Leu Leu Ly8 Met Arg Arg 35 40 45 Arg Arg Gin Phe Glu Phe Ser Val Asp Ser Phe Gln Ile Lys Leu Asp 35Ser Asn Asn Ser Gly Lys Asn Val Glu Leu Lys Phe Val Asp Ser Leu 20 30 Leu Glu Cys (2) INFORMATION FOR SEQ ID NO: 415: Glu Ile Arg Gly Gly Leu Leu Lys Tyr Cys Asn Leu Leu Val Arg 100 105 Ser Leu Leu Leu Phe Tyr Glu Cys Ser Glu Asn Pro Met Thr Glu Thr 50 55 (2) INFORMATION FOR SEQ ID NO: 416: Ala Phe Ξ Ξ (X (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 416: Asp His Leu Cys Asn Lys Ile Ile Ala Thr Arg Asn Pro Glu 85 90 95 SEQUENCE CHARACTERISTICS SEQUENCE CHARACTERISTICS: Ser Val Leu Phe Leu Pro Ile Ser Leu Asn Leu Leu Leu 20 30 Thr Ile Ile Gly Glu Ser Val Tyr Gly Asp Phe Gln Glu
70 75 80 SEQUENCE DESCRIPTION: SEQ ID NO: 415: (B) TYPE: amino acid
(D) TOPOLOGY: linear (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 32 amino acids (A) LENGTH: 163 amino acids

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Gly Phe Arg Pro Ala Ser Asp Glu Ile Lys Thr Leu Gln Arg Tyr Met

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2)

INFORMATION FOR SEQ ID NO:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 50 amino acids

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3 50 6 ઝ 30 25 20 2 5 Glu Glu Asn Arg Glu Thr Leu Lys Phe Tyr Leu Arg Ile Ile Leu Gly
20 25 30 Trp Phe Thr Ala Asp Ser Gly Thr Pro Ala Pro Glu His Asn Glu Lys 145 150 150 Ala Pro Gly Arg Ala Leu Tyr Leu Leu Trp Val Asn Val Leu Gly Pro 130 Gln Val Leu Ser Cys Phe Ser Leu Tyr Val Trp Ser Phe Trp Leu Leu 115 Gly Met Ala Glu His Leu Lys Asp Val Ile Leu Leu Thr Ala Ile Val 100' 105 110 Glu Asp Gly Ala Leu Met Asp Gly Gly Met Asp Leu Asn Met Glu Gln
95 Gly Ala Ser Tyr His Ser Met Ser Ser Met Ala Arg Ala Ala Phe Ser 65 70 75 80 Ala Ser Phe Trp Ala Trp Leu Ala Leu Gly Phe Ser Leu Ala Val Tyr 50 55 Ala Asn Ala Ile Tyr Cys Leu Val Thr Leu Val Phe Phe Tyr Ser Ser 35 40 45 Met Ala Pro Lys Gly Lys Val Gly Thr Arg Gly Lys Lys Gln Ile Phe 1 15 Gln Val Lys Leu Glu Ser Tyr Leu Gln Asn His Phe Val Gly Ile Gly Arg Pro 145 150 155 ş Arg Gln (2) INFORMATION FOR SEQ ID NO: 417: Ser Arg Phe Phe Ile Asp Phe Ser Asp Ile Gly Glu Gln Gln Arg 130 140 (i) SEQUENCE CHARACTERISTICS: Arg Arg Gln Glu Arg Arg Gln Met Lys Arg Leu Xaa 165 170 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 417: 15 (A) LENGTH: 174 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear 120 125

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 418: (B) TYPE: amino acid (D) TOPOLOGY: linear

Met Glu Leu Pro Lys Gly Leu Gln Gly Val Gly Pro Val Ala Met Met 1 15 1

Leu Pro Val Leu Cys Thr Gln Ala Leu Arg 25 ž 7yr Leu 1 20 Arg Pro Phe

Ser Gln Gly Lys Ser Pro Leu Leu Trp Lys Arg Thr Cys Cys Leu 35 40 듾

2

Ala Xaa 50

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(2) INFORMATION FOR SEQ ID NO: 419:

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SEQUENCE CHARACTERISTICS:
(A) LENGTH: 120 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 419:

22

Met Leu Gly Lys Gly Gly Gly Arg Ala Gly Leu Leu Arg Tyr Arg Leu  $_{\rm 1}$   $_{\rm 1}$ 

Leu Tyr Phe Thr Leu Val Val Gly Glu Gly Glu Pro Gly Glu Aan Lys 20 3

Thr Gly Lys Lys Ile Ile Phe Cys Ser 40 Phe Glu Thr Ile Pro Phe 35 35

Ser Asn Val Pro Ser His Lys Gly Pro Val 55 60 Lys Met Val Glu Asn

Leu Gly 80 Phe Leu Leu Ile Gly Arg Cys Ser Ser Gly Xaa 90 Trp Glu Leu Lys Ile Ser Glu Thr 75 Pro Leu Arg Ser Glu Gln 70 Glu Gly Lys Ile Gly 85 6

Leu Cys Phe Cys Trp Asp Val Leu Cys Cys Met Tyr Ala Tyr 100 g 45

25 E Arg Ser Leu Leu Ser 115 Asp Met

S

INFORMATION FOR SEQ ID NO: 420 3

(A) LENGTH: 159 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS

22

SEQUENCE DESCRIPTION: SEQ ID NO: 420: <del>(</del><del>x</del>

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Pro Glu Val Thr Gln Gln Thr Ile Glu Leu Lys Glu Glu Cya Lys 50 60 Met Thr His Leu Leu Leu Thr Ala Thr Val Thr Pro Ser Glu Gln Asn Ser Ser Arg Glu Pro Gly Trp Glu Thr Ala Met Ala Lys Asp Ile Leu 20 Gly Glu Ala Gly Leu His Phe Asp Glu Leu Asn Lys Leu Arg Val Leu 15 Asp Phe Val Asp Lys Ile Gly Gln Phe Gln Lys Ile Val Gly Gly Lev 65 Asp Gln Leu Ala Lys Glu Ala Glu Asn Glu Lys Met 85 Lys Ala Ile Gly Ala Arg Agn Leu Leu Lys Ser Ile Ala Lys Gln Arg 110 Glu Ala Gln Gln Gln Leu Gln Ala Leu Ile Ala Glu Lys Lys Met 115 Gin Leu Giu Arg Tyr Arg Val Glu Tyr Glu Ala Leu Cys Lys Val Glu 130 130 Ala Glu Gln Asn Glu Phe Ile Asp Gln Phe Ile Phe Gln Lys Xaa 145 Ile Glu Leu Val Asp 2 2 15 23 8

(2) INFORMATION FOR SEQ ID NO: 421:

35

(A) LENGTH: 154 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 421: ž.

Val Pro Val Ala 45 Met Asn Val Gly Val Ala His Ser Glu Val Asn Pro Asn Thr Arg Val Met Asn Ser Arg Gly Met Trp Leu Thr Tyr Ala Leu Gly Val Gly Leu 25 Trp Thr Leu Thr Asn Ile Ile His Asn Leu Gly Met Tyr Val Phe Leu Lys Ala Trp Glu Gln Leu Asp Tyr Gly Val Gln Phe Thr 90 95 Thr Pro Asp Gln Gly Ser Ile Pro Phe Phe Ser Thr Pro Phe Glu 70 Leu His Ile Val Leu Leu His Ala Val Lys Gly Arg Leu Leu Thr His 53 S <del>수</del> 5 55

Ser Ser Arg Lys Phe Phe Thr 11e Ser Pro 11e 11e Leu Tyr Phe Leu 100

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Gln Lys Asp Pro Arg Ala Asn Pro Ser Ala Phe Leu 195 , 200	Ser Phe Thr Glu Ile Leu Gly Val Trp Leu Thr Tyr Arg Tyr Arg Asn 185 180	Tyr Ala Gly Glu Val Leu Arg Phe Val Gly Gly Ile Gly Leu Phe Phe 175	val Lys Ser Asp His Ser Cys Ser Pro Cys Ala Pro Ile Ile Gly Glu 145 150 150	45 Cys Gly Phe Arg Ser Val Asn Pro Asn Asp Thr Cys Leu Ala Ser Cys 130 135	Asn Asn Thr Ala Ser Ala Arg Asn Asp Ile Gln Arg Asn Leu Asn Cys 115	Leu Ala Leu Asn Gln Glu Gln Gln Gly Gln Leu Leu Glu Val Gly Trp 100 105 110	Ile Leu Leu Leu Val Phe Ile Val Gln Phe Ser Val Ser Cys Ala Cys 85	Ile Gly Ala Val Lys His His Gln Val Leu Leu Phe Phe Tyz Met Ile 65 70 75 80	00 Val Ile Ala Val Gly Ile Phe Leu Phe Leu Ile Ala Leu Val Gly Leu 50	Trp Gly Ile Gly Phe Gly Leu Ile Ser Ser Leu Arg Val Val Gly Val 35 40 45	Asn Leu Leu Tyz Thr Leu Val Ser Leu Leu Leu Ile Gly Ile Ala Ala 25 29	Met Val Cys Gly Gly Phe Ala Cys Ser Lys Asn Cys Leu Cys Ala Leu 1 15	5 (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 204 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 422:	(2) INFORMATION FOR SEQ ID NO: 422:	Val Arg Ile Phe Gly Ile Asn Lys Tyr Xaa 145 0	5 Ala Ser Leu Leu Ser Val Leu Île Pro Lys Met Pro Gln Leu His Gly 130 135	Ala Ser Phe Tyr Thr Lys Tyr Asp Pro Thr His Phe Ile Leu Asn Thr 115 120
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25 20 5 5 Tyr Met Xaa 65 Met Leu Gln Ser Ile Ile Lys Asn Ile Trp Ile Pro Met Lys Pro Tyr
1 10 15 Phe Ile Val Tyr Lys Ile Arg Ala Ala Asp Lys Arg Ser Lys Ala Leu 35 40 45 Lys Ala Ser Ala Pro Ala Pro Gly His His Asn Gln Ile Tyr Leu Glu 50 55 Tyr Thr Lys Val Tyr Gln Glu Ile Trp Ile Gly Met Gly Leu Met Gly 20 25 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 423:

(2) INFORMATION FOR SEQ ID NO: 424:

30 35 Met Leu Gly Val Ser Leu Pho Leu Leu Val Val Leu Tyr His Tyr Val 1 15 (i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 25 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 424:

(2) INFORMATION FOR SEQ ID NO: 425:

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Ala Val Asn Asn Pro Lys Lys Gln Glu 20 25

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45 50 S Gly Gly Ala Gly Ala Pro Ser Gly Thr Val Pro Val Leu Phe Cys Phe 20 25 30 Met Ala Ala Xaa Glu Pro Ala Val Leu Ala Leu Pro Asn Ser Gly Ala 1 15 (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 299 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 425:

Leu Leu Ile Gln Lys Phe Leu Ser Leu Tyr Gly Asp Gln Ile Asp Met

Ser Val Phe Ala Arg Pro Ser Ser Val Pro His Gly Ala Gly Tyr Glu 45

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(2) INFORMATION FOR SEQ ID NO: 423:

(i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 67 amino acids

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His Arg Lys Phe Val Val Gln Leu Phe Ala Glu Glu Trp Gly Gln Tyr 75

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Ser Glu Arg Cys Lys Val Arg 90 Leu Pro Lys Gly Phe Ala Val 85 Val Asp

Leu Gln Ile Gln Leu Thr Thr Leu Gly Asn Leu Thr Pro 100 100 Pro Leu Val 9 Ser Ser Thr Val Phe Phe Cys Asp Met Gln Glu Arg Phe Arg Pro 120

Ala Ile Lys Tyr Phe Gly Asp Ile Ile Ser Val Gly Gln Arg Leu Leu 130 2

Gln Gly Ala Arg Ile Leu Gly Ile Pro Val Ile Val Thr Glu Gln Tyr 145 Pro Lys Gly Leu Gly Ser Thr Val Gln Glu Ile Asp Leu Thr Gly Val 175 20

Lys Leu Val Leu Pro Lys Thr Lys Phe Ser Met Val Leu Pro Glu Val 180 25

Glu Ala Ala Leu Ala Glu Ile Pro Gly Val Arg Ser Val Val Leu Phe 195

Gly Val Glu Thr His Val Cys Ile Gln Gln Thr Ala Leu Glu Leu Val 210 8

Ser Ser Arg Met Met Asp Arg Met Phe Ala Leu Glu Arg Leu Ala Xaa Kaa Gly 250 255 Gly Arg Gly Val Glu Val His Ile Val Ala Asp Ala Thr 225 Ser

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The The See Glu Ala Val Leu Leu Gln Leu Val Ala Asp 260 Lys Asp His Pro Lys Phe Lys Glu Ile Gln Asn Leu Ile Lys Ala Ser 275 Ile Ile Val <del>6</del>

Ser Lys Val Xaa 295 295 Le Le Pro Glu Ser Gly 290 Ala 45

(2) INFORMATION FOR SEQ ID NO: 426: လ

(A) LENGTH: 13 amino acids SEQUENCE CHARACTERISTICS: TYPE: amino acid 3

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 426:

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Met Arg Asp Leu Gly Thr Leu Leu Ser Pro Val Cys Ser 1 5 S

INFORMATION FOR SEQ ID NO: 427; 3

(A) LENGTH: 198 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear

427: SEQUENCE DESCRIPTION: SEQ ID NO: ź.

Phe Gly Cys Leu Val Ala Gly Arg Leu Val Gln Thr Ala Ala Gin 15 Ser Gln Val Ala Glu Asp Lys Phe Val Phe Asp Leu Pro Asp Tyr Glu  $25\ \ \,$  30 Met 9 15

ile Asn His Val Val Val Phe Met Leu Gly Thr Ile Pro Phe Pro Glu 35

Met Gly Gly Ser Val Tyr Phe Ser Tyr Pro Asp Ser Asn Gly Met 50 ξ

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Lys Pro Ser Ala Pro Val Trp Gln Leu Leu Gly Phe Val Thr Asn Gly 65

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Pro Pro Ser Val Ala Gln Ile Gly 110 lle Phe Lys Ile Ser Gly Leu Lys Ser Gly Glu Gly Ser Gln His 95 7 Phe Gly Ala Met Asn Ile Val Arg

8

ile Ser Val Glu Leu Leu Asp Ser Met Ala Gln Gln Thr Pro Val Gly 115

Phe Thr Gln Phe Thr Gln Lys 140 Ser Asn Ala Ala Val Ser Ser Val Asp 130 35

Met Leu Asp Asn Phe Tyr Asn Phe Ala Ser Ser Phe Ala Val Ser Gin 145 Ala Gin Met Thr Pro Ser Pro Ser Giu Met Phe Ile Pro Ala Asn Val 175 5

Val Leu Lys Trp Tyr Glu Asn Phe Gln Arg Arg Leu Ala Gln Asn Pro 190

Xaa Phe Trp Xaa Thr Xaa 195

45

(2) INFORMATION FOR SEQ ID NO: 428:

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(A) LENGTH: 47 amino acids SEQUENCE CHARACTERISTICS (B) TYPE: amino acid 3

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 428:

Met Gly Leu Pro Leu Met Ala Leu Met Trp Ser Thr Leu Pro Ala Ser 8

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S Ala Gly Val Asn Phe Ile Leu Ala Leu Pro Leu Leu Leu Leu Trp Lys 20Ile Lys His Trp Ile Thr Ile Ile Arg Ala Arg Phe Glu Glu Val Leu \$45\$Met Gly Glu Val Ile Leu Ala Val Cys His Pro Asp Cys Ile Thr Thr 20Met Lys Lys Val Glu Glu Lys Arg Val Asp Val Asn Ser Ala Val 1 5 10 Asn Arg Gly Gly Val Gly Arg Ser Val Met Ser Ala Val Glu Xaa 35 40 Glu Glu Met Thr Arg Lys Gln Pro Asp Val Asp Arg Val Thr Lys Thr 115 120 125 끍 (2) INFORMATION FOR SEQ ID NO: 429: Asn Ile Asp Arg Val Lys Ala Leu Ile Ala Glu His Gln Thr Phe Met  $100\,$ Trp Ala Glu Thr Thr Leu Ile Gln Arg Asp Gln Glu Pro Ile Pro Gln 95 Leu Val Ala Asn Ala Glu Leu Leu Glu Glu Leu Leu Ala Trp Ile Gln 65 70 75 Ser Arg Ser Gly Gly Arg Lys Ser Leu Ser Gln Pro Thr Pro Pro Pro 145 Tyr Lys Arg Lys Asn Ile Glu Pro Thr His Ala Pro Phe Ile Glu Lys 130 140 Ala Asn Phe Asp Phe Asp Val Trp Arg Lys Lys Tyr Met Arg Trp Met 210  $^{\circ}$  220  $^{\circ}$ ğ Leu Ser Ala Arg Trp Gln Gln Val Trp Leu Leu Ala Leu Glu Arg Gln . 180 185 Met Pro Trp Ala Lys Gln His Gln Gln Arg Leu Glu Thr Ala Leu Ser Glu 50 56 Lys Leu Asn Asp Ala Leu Asp Arg Leu Glu Glu Leu Lys Glu Phe 195 200 205 ξ (1) SEQUENCE CHARACTERISTICS: Ile Leu Ser Gln Ser Glu Ala Lys Asn Pro Arg Ile Asn Gln 165 170 175 SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 370 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear 5 25 Ala

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ઝ 25 20 2 5 Gly Xaa 370 Ile Phe Asp Arg Asp Gly Asp Gly Tyr Ile Asp Tyr Tyr Glu Phe Val 275Ala Ser Lys Phe Pro Thr Thr 260 Asp Gln Asp Gly Lys Ile Thr Arg Gln Glu Phe Ile Asp Gly Ile Leu 245 250 256 Asn His Lys Lys Ser Arg Val Met Asp Phe Phe Arg Arg Ile Asp Lys 225 230 230 Ile Leu Arg Asn Arg Asp Gly Ser Arg Trp Trp Arg Met Asp Gly Leu 355 Phe Leu Gly Asn Gln Phe Gly Asp Ser Gln Gln Leu Arg Leu Val Arg 340 345 Ala Ala Leu His Pro Asn Lys Asp Ala Tyr Arg Pro Thr Thr Asp Ala 290 295 Ala Lys Arg Asp Lys Ile Glu Asp Glu Val Thr Arg Gln Val Ala Gln Cys Lys Cys 305 310 2 INFORMATION FOR SEQ ID NO: 430: Phe Gin Val Glu Gin Ile Gly Glu Asn Lys Tyr Arg Phe 325 330 335 Lys Leu Glu Met Thr Ala Val Ala Asp 265 270

(2) INFORMATION FOR SEQ ID NO: 431:

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Leu Tyr Leu Arg Tyr Val Thr Phe Val Tyr Leu Asn Leu Phe
20 25 30

Met Asn Val Lys Thr Phe Ser Xaa Asp His Met His Phe Leu Cys Cys
1 10 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 430:

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 30 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

E ξ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 431: (A) LENGTH: 24 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Met Glu Pro His Leu Arg Cys Arg Val Thr Arg Val Arg Gly Ser Leu 1 15

8 Gly Asn Thr Gly Arg Trp Leu Leu

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		614
	20	Val Trp Ser Pro Ser Thr Ser Arg Leu Thr Arg Tyr Thr 11e Trp His 145
2	(2) INFORMATION FOR SEQ ID NO: 432:	5 Leu Gln Pro Pro Leu Gln Thr Thr Cys Ile Ile Leu Ser Arg His Xaa
10	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 53 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 412:	01
	Met His Tyr Leu Val Leu Gly Gly Leu Gly Val Phe Leu Phe Ser $_1$ $_1$ $_2$ $_3$ $_4$ $_1$ $_2$ $_1$ $_2$ $_3$ $_4$ $_2$ $_3$ $_4$ $_4$ $_5$ $_4$ $_5$ $_4$ $_5$ $_5$ $_5$ $_7$ $_7$ $_7$ $_7$ $_8$ $_7$ $_8$ $_7$ $_8$ $_9$ $_9$ $_9$ $_9$ $_9$ $_9$ $_9$ $_9$	(2) INFORMATION FOR SEQ ID NO: 434:
15	Phe Ala Phe Phe E	5 (1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 77 amino acids (B) TYPE: amino acid (D) TOPOLOGY: 1lnear
70	Tyr Leu Glu Gly Met Gly Gly Ser Gly Asn Arg Glu Val Gly Gly Gly Jy 15 40 45	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 434:
	Phe Cys Leu Phe Phe 50	net bed Arg Cys inp tto bed rise inp bed to bed out to rise in
25		Cys Ser Leu Phe Trp Leu Leu Val Glu Trp Phe Gly Thr Asn Ile Asp 25 25 20 20 20 20 25 25 25 25 25 25 25 25 25 25 25 25 25
	(2) INFORMATION FOR SEQ ID NO: 433:	Arg Glu Ser Tyr Asp Ala Ile Gly Gly Pro Ser Trp Met Thr Ala Ser
30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 176 amino acids (B) TYPE: anino acid (D) TOPOLOGY: linear	30 Ser Phe Cys Leu Ser Asn Ser Asn 11e Trp Ser Leu Glu I1e Ser Ser 50 55
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 433: Met Val Ser Lys Ala Leu Leu Arg Leu Val Ser Ala Val Asn Arg Arg 1 5 1 10	Gly Ser Thr Ser Val Val His Ser Gin Gin Ala Met Asp 65 75 35
	Arg Met Lys Leu Leu Leu Gly Ile Ala Leu Leu Ala Tyr Val Ala Ser 20	(2) INFORMATION FOR SEQ ID NO: 415:
40	-	40 (i) SEQUENCE CHARACTERISTICS: (a) LENGTH: 32 and no acids (b) TYPE: and no acid (c) TOPOLOGY: linear
45	Leu Lys Ile Glu Ser Lys Ile Glu Glu Met Val Glu Pro Leu Arg Glu 50 50	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 435:
	Lys lie Arg Asp Leu Glu Lys Ser Phe Thr Gln Lys Tyr Pro Pro Val 65 80	Met Arg Ser Cys Glu 11e Gln Leu Cys Val Trp Leu Leu Val Ser Ser 1 5 10 10 115 115 115 115 115 115 115 1
20	Lys Phe Leu Ser Glu Lys Asp Arg Lys Arg Ile Leu Ile Thr Gly Gly 85	Als Val Asp Met Val Leu Gly Gly Ser Fro Ser 'nn Leu lyr Met Met 30 20 25 30 30 30 30 30 30 30 30 30 30 30 30 30
55		
	nts ctu vat vat vat kep fele ille ut cty Aky Lys Aky Ash 125 125 125 125 125 125 125 125 125 125	(2) INPORMATION FOR SEQ ID NO: 436:
9	Val Glu His Try Ile Gly His Glu Asn Phe Glu Leu Ile Asn His Asp 130	(1) SEQUENCE CHARACTERASILLOS: (A) LENGTH: 30 amino ecids

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Leu Pro Cys Thr Pro Gln Met Val Arg Gly Val Thr Gln Val Leu Arg  $50 \ \ \, 55$ Glu Leu Ser Thr Asp Ser Ser Ala Arg Leu Leu Tyr His Glu
20 25 30 Met Val Val Asn Ser Leu Cys Phe Leu Ser Leu Leu Leu Val Ile Leu 1 10 15 Glu Phe Gly Asp Gln 65 Gln Gly Lys Ile Ala Phe Sex Leu Met Phe Val Leu Lys Asp Leu Sex 20 25 30 Met Asp Lys Gln Lys His Leu Glu Val Arg Arg Ser Val Phe Lys Ile 1 10 Pro Thr (2) INFORMATION FOR SEQ ID NO: 437: <u>E</u> (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 436: Tle Phe Ser His Ser Ile Leu Leu Leu Leu Pro His His Val 35 40 45 SEQUENCE DESCRIPTION: SEQ ID NO: 437: (B) TYPE; amino acid
(D) TOPOLOGY: linear (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 69 amino acids 615

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(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 440:

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Arg Val Met Val Asn Leu Asn Ile Leu Phe Xaa 35 40

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Leu Leu

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Phe Leu 20

Phe

Pro Leu Ser Phe Leu Pro Asn Ile Trp Arg 25

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33 30 ટ્ડ 20 45 8 8 55 5 Leu Thr Thr Leu Leu 50 Feu Met Trp Gln Val Arg Gly Leu Pro Pro Val Pro Leu Leu Leu Thr Met
1 15 2 Asn Pro Ser Arg Leu Ile Leu Tyr Met Ile Ser Ala Gly Ala Asp Ser 20 25 30 Met Leu Leu Phe Pro Ser Leu Leu Phe Ala Ala Thr Tyr Asn Val Ala 1 15 3 Phe Glu Ala Val Pro Ile Ser Val Ser Asp Gln Pro Ser Pro Xaa 35 40 45 Pro Pro Pro Cys Leu Ser Ser Pro Phe Pro Phe Ile Ser Val Pro  $20 \ 25 \ 30$ INFORMATION FOR SEQ ID NO: 441: INFORMATION FOR SEQ ID NO: 442: (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 440: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 441: (B) TYPE: amino acid
(D) TOPOLOGY: linear (D) TOPOLOGY: linear (A) LENGTH: 33 amino acids (B) TYPE: amino acid (A) LENGTH: 53 amino acids

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Met Lys Phe Ser Leu Val Leu Lau Ila Lys Ila Ila Ser Pha Glu Arg 1 5 10 15

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(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 442:

(B) TYPE: amino acid (A) LENGTH: 64 amino acids

(i) SEQUENCE CHARACTERISTICS:

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 439: (D) TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 43 amino acids(B) TYPE: amino acid

(2) INFORMATION FOR SEQ ID NO: 439:

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SEQUENCE DESCRIPTION: SEQ ID NO: 438:

(D) TOPOLOGY: linear

Met Pro Leu Cys Phe Phe Ser Phe Leu Cys Cys Trp Val Leu Val Phe 1 19

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(2) INFORMATION FOR SEQ ID NO: 438:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 19 amino acids
(B) TYPE: amino acid

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Lau Ile Phe Phe Phe Pro Tyr Leu Ser Leu Val Thr Leu Leu Gln Ala Arg Asn Leu Trp Val 11e His Arg Ala Ala Leu Cys Glu Ser Gly Leu Phe His Trp Arg Lys Gly Ila Glu Asn Gln Leu Glu Pro Met Tyr Phe Leu Pro His Gly Thr Leu Phe Leu Met Ile Thr Ser Val Leu Val Phe 9

(2) INFORMATION FOR SEQ ID NO: 443: 8

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Leu Tyr Ser Cys Glu Pro Tyr Leu Ile Ile Leu Asn Ile Tyr Ser 5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 443: SEQUENCE CHARACTERISTICS:
(A) LENGTH: 34 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear Met 22

Gln Lys Ala Phe Tyr Phe Tyr Phe Bhe Glu Gly Ser Phe Ser Val Cys  $$20\ \ \, 20$ 3

Thr Leu

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(2) INFORMATION FOR SEQ ID NO: 444: 9

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 444: SEQUENCE CHARACTERISTICS:
(A) LENGTH: 89 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Leu Ala Gln Glu Cys Pro Pro His Ile Pro Ser Ser Phe Phe Leu Val 20 Met Arg Gln Arg Gln Ala Ala Cys Gln Pro Pro Pro Ser Arg Asn Gly

Leu Leu Phe Ile Pro Trp Leu Ala Ser Leu Leu Ser Ser Pro Leu 35 45 ξ တ္တ

Leu Gln Gln Cys Arg Gln His Gln Val Leu Gln Glu Lys Asn Thr  $$70\ \ \, 70\ \ \, 75\ \ \, 80$ Ser Ile Ser Trp Asp Leu Gly Leu Lys Leu 55 60 Leu Val ž Asn Leu Leu I 50 Agn 65 55

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Lys Lys Phe Asn Lys Lys Lys Lys Lys Lys 85

(2) INFORMATION FOR SEQ ID NO: 445:

(i) SDQUENCE CHARACTERISTICS:
(A) LENGTH: 350 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 445:

Met Asp Phe Ile Thr Ser Thr Ala Ile Leu Pro Leu Leu Phe Gly Cys 1 5 10 15 Phe Gly Leu Phe Arg Leu Leu Gln Trp Val Arg Gly Lys  $20\ \ 25$ 

Leu Gly Val

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Leu Gly Lys Glu Cys Ala Lys Val Phe Tyr Ala Ala Gly Ala Lys Leu 50 Arg Asn Ala Val Val Ile Thr Gly Ala Thr Ser Gly 35 55 Ala Tyr

Val Leu Cys Gly Arg Asn Gly Gly Ala Leu Glu Glu Leu Ile Arg Glu 65 75 80

22

Leu Thr Ala Ser His Ala Thr Lys Val Gln Thr His Lys Pro Tyr Leu 8 Glu Ile Leu Gln Cys Phe Gly Tyr Val Asp Ile Leu Val Asn Asn Ala 115 33

Val Thr Phe Asp Leu Thr Asp Ser Gly Ala Ile Val Ala Ala Ala Ala 100

Leu Thr Lys 160 Gly Ile Ser Tyr Arg Gly Thr Ile Met Asp Thr Thr Val Asp Val Asp 130 Lys Arg Val Met Glu Thr Asn Tyr Phe Gly Pro Val Ala 145 155 145 6

Ala Leu Leu Pro Ser Met Ile Lys Arg Arg Gln Gly His Ile Val Ala 175 45

Ile Ser Ser Ile Gin Gly Lys Met Ser Ile Pro Phe Arg Ser Ala Tyr 180

Ala Ala Ser Lys His Ala Thr Gln Ala Phe Phe Asp Cys Leu Arg Ala 195 ဂ္ဂ

Glu Met Glu Gln Tyr Glu Ile Glu Val Thr Val Ile Ser Pro Gly Tyr 210 Ile His Thr Asn Leu Ser Val Asn Ala Ile Thr Ala Asp Gly Ser Arg 225

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Tyr Gly Val Met Asp Thr Thr Thr Ala Gln Gly Arg Ser Pro Val Glu 255

Val Ala Gln Asp Val Leu Ala Ala Val Gly Lys Lys Lys Lys Asp Val 260 265

S 55 Met Ala Sar Ala Glu Leu Asp Tyr Thr Ile Glu Ile Pro Asp Gln Pro
1 5 10 15 Trp Ser Gln Lys Asn Ser Pro Ser Pro Gly Gly Lys Glu Ala Glu
20 25 30 Arg Gln Pro Val Val Ile Leu Leu Gly Trp Gly Gly Cys Lys Asp 35  $^\dagger$  40  $^45$ (1) SEQUENCE CHARACTERISTICS: (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 447: (A) LENGTH: 278 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

6 **A**gn 2 Leu Trp Leu INFORMATION FOR SEQ ID NO: 446: E £ SEQUENCE CHARACTERISTICS: Thr Phe Ile Leu Glu Thr Glu Val Tyr Leu 25 SEQUENCE DESCRIPTION: SEQ ID NO: 446: (A) LENGTH: 49 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

50 3 5 INFORMATION FOR SEQ ID NO: 447:

Glu Ala Ala Leu Leu Gly Leu Leu Thr Leu Gln Gly Thr Val Ala Phe 325 330 335 Gly Asn Pro Arg Thr Pro Ser Thr Leu Thr Ser Gln Gly Gln Gly Arg 305 310 315 Ile Leu Ala Asp Leu Leu Pro Ser Leu Ala Val Tyr Leu Arg Thr Leu 275 280 285 Pro Gly Leu Phe 290 Glu Thr Leu Met Glu Ile Cys Leu Thr Ser Gly Lys Asp 345 350 Phe Ser Leu Met Pro Pro Gly Pro Glu Lys Ser 295

20

Ile Leu Glu Arg Arg Ala Ala Met Leu Arg Leu Leu Leu Leu Val Ala 165 170 175

Phe Ala Leu Val Val Leu Phe His Val Leu Leu Ala Pro Ile Thr 180 185

Ala Pro Gly Asp Ser Asn Leu Val Gly Ala Leu Arg Ala Leu Ala Ala 145 150 150

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Ser Asn

Gly Gly Val Met Leu 115

Tyr Arg Tyr Val Leu Glu Leu Leu Gln 120 125

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Arg Arg Phe Cys Arg Leu Arg Val Val Oly Thr Ile Phe Asp Ser 130 140

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Leu Gly Ile Pro

Ser Leu Arg Val Leu 85

Ala Gln Lys Leu Leu Glu Leu 90 95

Leu Phe Asp Tyr Glu Ile Glu Lys Glu Pro Leu Leu Phe His Val Phe
100 105

Met Val Phe Leu Pro Arg Gly Val Val Val Ser Gly Gly Ala Ala Cys 1 5 10 15

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Arg Trp Pro Glu Leu Tyr Leu Tyr Ser Arg Ala Asp Olu Val Val Leu 210 \$210\$

Ala Arg Asp Ile Glu Arg Met Val Glu Ala Arg Leu Ala Arg Arg Val 225 230 230

Ala Xaa Phe His Thr His Phe Tyr Asp Arg Leu Oln Asp Ala Gly Ser 195 200 205

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Ala

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1 Asp Leu Ala 30

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Thr Glu Ala Arg Ala His Ser Arg Met Gly Leu Gly Leu Trp Pro Pro 35  $$40^{\circ}$$ 

6 Leu Ala Arg Ser Val Asp Phe Val Ser Ser Ala His Val Ser His Leu 245 255

Arg Asp Tyr Pro Thr Tyr Tyr Thr 260 Ser Leu Cys Val Asp Phe Met Arg 265 270

Asn Cys Val Arg Cys Xaa 275

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(2) INFORMATION FOR SEQ ID NO: 448:

 $\widehat{\Xi}$ SEQUENCE CHARACTERISTICS (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 199 amino acids

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Met Ser Phe Ile Phe Asp Trp Ile Tyr Ser Gly Phe Ser Ser Val Leu 1 15

Cln Phe Leu Cly Leu Tyr Lys Lys Thr Cly Lys Leu Val Phe Leu Gly
20 25 30

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SEQUENCE DESCRIPTION: SEQ ID NO: 448:

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Lys Asn Leu Ala Lys Tyr Ser Ala Ile Tyr His Lys Arg Gly Cys Ile 50 60

Val Ile Arg Tyr Thr Ala Pro Trp His Met Val Phe Phe Ser Glu Ser 65 70 75 80

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Leu Asp Asn Ala Gly Lys Thr Thr Leu Leu His Met Leu Lys Asp Asp 35

Arg Leu Gly Gln His Val Pro Thr Leu His Pro Thr Ser Glu Glu Leu 50

Thr lle Ala Gly Met Thr Phe Thr Thr Phe Asp Leu Gly Gly His Val 65 75 80 으

Gin Ala Arg Arg Val Trp Lys Asn Tyr Leu Pro Ala Ile Asn Gly Ile 95

Ser Lys Val Phe Leu Val Asp Cys Ala Asp His Glu Arg Leu Leu Glu 100 105 2

Glu Glu Leu Asp Ser Leu Met Thr Asp Glu Thr Ile Ala Asn Val Pro 115

Leu lle Leu Gly Asn Lys lle Asp Arg Pro Glu Ala lle Ser Glu 130 116 2

Glu Arg Leu Arg Glu Met Phe Gly Leu Tyz Gly Gln Thr Thr Gly Lys 145

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Gly Ser Ile Ser Leu Lys Glu Leu Asn Ala Arg Pro Leu Glu Val Phe 175 Ser Val Leu Lys Arg Gln Gly Tyr Gly Glu Gly Phe Arg Trp 180 Met Cys 39

Met Ala Gln Tyr Ile Asp Xee 195

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(2) INFORMATION FOR SEQ ID NO: 449:

(B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 449:

Met Thr Leu Ser Arg Phe Ala Tyr Asn Gly Lys Arg Cys Pro Ser Ser 45

Tyr Asn Ile Leu Asp Asn Ser Lys Ile Ile Ser Glu Glu Cys Arg Lys 20

絽 Glu Leu Thr Ala Leu Leu His His Tyr Tyr Pro Ile Glu Ile Asp 35 40 8

Lys Ala His Asn Leu Leu Cys Gln Gln Lys Ile Gln Lys Phe Gln Ile 65 75 80 His Arg Thr Val Lys Glu Lys Leu Pro His Met Val Glu Trp Trp Thr 50 25

Ala Gln Val Val Arg Glu Ser Asn Ala Met Leu Arg Glu Gly Tyr Lys 8

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95 8 82 Thr Phe Phe Asn Thr Leu Tyr His Asn Asn Ile Pro Leu Phe Ile Phe 100 Ser Ala Gly Ile Gly Asp Ile Leu Glu Glu Ile Ile Arg Gln Met Lys 115

Phe His Pro Asn Ile His Ile Val Ser Asn Tyr Met Asp Phe Asn 130 Val

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Phe Leu Gln Gly Phe Lys Gly Gln Leu Ile His Thr Tyr 150 150 Glu Asp Gly 1 145 Ash Lys Ash Ser Ser Val Cys Glu Ash Xaa Gly Tyr Phe Gln Gln Leu 165 Thr Asn Val Ile Leu Leu Gly Asp Ser Ile Gly Asp Leu 180 180 Glu Gly Lys

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Thr Met Ala Asp Gly Val Pro Gly Val Gln Asn Ile Leu Lys Ile Gly 105 195

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Leu Asn Asp Lys Val Glu Glu Arg Arg Xaa Arg Tyr Met Asp Ser 210 Tyr Asp Ile Val Leu Glu Lys Asp Glu Thr Leu Asp Val Val Asn Gly 225 Phe 22

ile Leu Cys Gln Gly Val Gln Leu Glu Met Gln Gly 245 Leu Leu Gln His ೫

Pro Xaa

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INFORMATION FOR SEQ ID NO: 450: 3

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SEQUENCE DESCRIPTION: SEQ ID NO: 450;

Ser His Val Leu Leu Cys Pro Ser Leu Ser Cys Ser Asn Leu Leu 18 10 10 Leu Ser Pro His Leu 30 Pro Pro Ser His Ser Leu Gly Thr Met Gly Ser 20 25 45 င္လ

Ser Pro Thr Leu Cys Gly His Thr Met Cys Pro Val Asn Pro Glu Leu Pro Leu Ser 15 Ser Thr Thr Asp Gln Pro Gln Pro Asp Ala Cys 55 60 55

Leu Thr Leu Pro Leu Pro Ser Ser Phe Leu Pro His Ser Lys Pro Thr 65 75 80

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Gly Ala Ile Gly Lys Val Tyr His Ala Leu Asn Pro Lys Leu Thr Val 245 250 250 Ala Asn Thr Lys Tyr Ala Gin Asp Tyr Asn Pro 235 230 230 Phe Trp Cys Tyr Lys 240

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Gly Val Phe Pro Val Thr Val Gln Pro Pro His Cys Val Pro Asp Thr 195 200 205 볶 Ser Asn Ala Thr Leu Trp Tyr Lys Ile Phe Thr Thr Ala Arg Asp 210 215

Cys Glu Gln Val Val Phe Thr Ala Cys Met Thr Leu Thr Ala Ser Pro 180 185

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Lys Val Tyr Gln Lys Thr Pro Ala Leu Ala Val Lys Ala 40 45

Gly Phe Ile Ser Gly Trp Asn Leu Val Ser Met Cys Val Glu Tyr Val  $25 \ \ 20$ 

Met Ala Val Gly Gln Ile Met Thr Phe Gly Ser Pro Val Ile Gly Cys
1 10 15

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SEQUENCE DESCRIPTION: SEQ ID NO:

(A) LENGTH: 383 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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INFORMATION FOR SEQ ID NO: 453:

(1) SEQUENCE CHARACTERISTICS:

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Thr Leu Pro Thr Ala Trp Ser Ser Asp 165 Asp Cys Ala Leu His Gly His 170

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Ile Gly Leu Ser Gly Arg Glu Ala His Glu Glu Ile Asn Ile Thr Phe 145 150 150

볶 Ser Arg Asn Val Thr His Leu Tyr Ser Thr Ile Leu Gly His Gln 130 140

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Ser Val Ser Ile Thr Leu Thr Leu Asp Pro Leu Lys Pro Phe Gly Gly 115 120 128

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Ser The Gln Ser Pro Gln Ala Leu Glu Asp Ser Gly Pro Val Asn Ile 100 105

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Asp Thr Thr 쿥 Pro Glu Ser Thr Met 85 Thr Ser Gly Gln Ala Arg Ala 90 95

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Val Asn Gly Glu Ala Ala Leu Ser Pro Gly Leu Cys Asp Pro Ile Ser . 35 40 45

Glu Ala Leu His Ala Gly Cys Phe Pro Ala Phe Ala Ser Ala Thr Arg 20 25

Met Pro Gly Leu Ser Leu Ala Leu Leu Pro Phe Gly Pro Gly Cys Thr 1 15

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Val Pro Tyr Val 50

Leu Asp Leu Cys Val Ser Glu Asn Glu Thr Leu Lys His Leu Thr Asn 65 70 75 80

Pro Glu Met Ala Glu Asp Trp Asn Thr Phe Leu 50 99 19 Arg Phe Asn Αsp

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Ala Phe Leu Thr Leu Gly Tyr Phe Phe Lys Ile Lys Glu Ile Lys Ser 35

 $\mbox{Arg Pro Pro Leu Val Val Phe Met Ile Ser Val Xaa Pro Met Ala Ile 20 <math display="inline">25$  30

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Xaa His Pro Cys Ser Pro 85

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SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 315 amino acids INFORMATION FOR SEQ ID NO: 451:

Met Phe Ser Ile Asn Pro Leu Glu Asn Leu Lys Val Tyr Ile Ser Ser 1 15

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SEQUENCE DESCRIPTION: SEQ ID NO: 451:

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(2) INFORMATION FOR SEQ ID NO: 452:

(1) SEQUENCE CHARACTERISTICS

£ SEQUENCE DESCRIPTION: SEQ ID NO: 452: (A) LENGTH: 52 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Cys Pro Glu I 305 Lys Val Ala Leu Ala Glu Ala Xaa 310 315

Š 쿭 Ile Lys Gly Arg Pro Ser Lys Leu Arg Gln Ser Asn Pro Glu Phe 290 300 Ser Tyr Phe Leu l 275 Phe Val Met Val Ile Thr Met Phe Cye Tyr Ala 280 285 Ile Val Pro Asp Asp Asp Arg Ser Leu Ile Asn Leu His Leu Met His 260 265

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G) Y

Leu Lys Glu Glu Glu Thr Glu Leu Lys Gln Leu Asn Leu His Lys 50 60

Asp Thr Glu Pro Lys Pro Leu Glu Gly Thr His Leu Met Gly Val Lys 65 70 75 80

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Thr Ser 175 Asp Ser Asn Ile His Glu Leu Glu His Glu Glu Glu Pro Thr Cys Ala 95 Phe Arg Asp Gly Trp Val Ser 110 Leu Ala Phe Leu 125 Thr Gln Cly Leu Ser Gly Phe His Pro Gln Tyr Phe Asp Gly Ser Ile 145 Phe Asp Cys Ile Thr Thr Gly Tyr Ala Tyr 135 Lys Met Trp Phe Gly Ser Ala Gly Leu Ile Ser Gly Leu Ala Gln Leu 180 Ser Cys Leu Ile Leu Cys Val Ile Ser Val Phe Met Pro Gly Ser Pro 195 Phe Ile Gin Gly Glu Ser Ile Thr Pro Thr Lys Ile Pro Glu Ile Thr Thr Glu 225 Pro Glu Ser Val Pro Ile Ile Ser Val Ser Leu Leu Phe Ala Gly 260 Leu Trp Ser Phe Asp Leu Thr Val Thr 280 Met Val Ile Leu Ala Pro Asn Pro Glu Ala Phe Gly Leu Leu Val Leu 335 Lys Leu Phe Ala Cys Gly Pro Asp Ala 360 Ile Tyr Met Ser Asn Gly Ser Asn Ser Ala Asn Ile Val Pro Glu Thr 255 Leu Leu Gln Glu Asn Val Ile Glu Ser Glu Arg Gly Ile Ile Asn 290 Phe Ile Ser Phe Val Ala Met Gly His Ile Met Tyr Phe Arg Phe 340 : Ser Val Val Xaa 380 Ser Tyr Asn Trp Asn Asn Gly Asn Cys Ser Phe Tyr Leu Ala 165 Leu Asp Leu Ser Val Ser Pro Phe Glu Asp Ile Arg Ser Arg 210 Val Gln Asn Ser Met Asn Tyr Leu Leu Asp Leu Leu His 316 Tyr Tyr Asn Gln Pro Val Phe Leu Aia Gly Met Gly 115 Asn Thr Ser Gln Met Ala Glu Pro Phe Arg Thr 100 Glu Val Arg Lys Glu Asn Gln Ala 370 370 Ą Val Ile Ala Ala Arg Ile Gly Thr Val Leu Gly Ala Gln Asn Thr Leu Gly 355 Ser Val Met 130 꿏 G1y 305 Ser 당 116 Lya 2 2 15 22 8 35 6 တ္တ 22 5

(2) INFORMATION FOR SEQ ID NO: 454:

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acida (A) LENGTH: 186 amino (i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 454:

Met Arg Ser Ile Gly Asn Lys Asn Thr Ile Leu Leu Gly Leu Gly Phe 1 5

Met Trp Ala Ala Gly Ala Val Ala Ala Met Ser Ser Ile Thr Phe Pro 35 Gin ile Leu Gin Leu Ala Trp Tyr Gly Phe Gly Ser Glu Pro Trp Met 2

Ala Val Ser Ala Leu Val Ser Arg Thr Ala Asp Ala Asp Gln Gln Gly 50 60 15

2 g Gly Pro Ala Leu Tyr Gly Phe Ile Phe Tyr Ile Phe His Val Glu Leu Val Val Gln Gly Met Ile Thr Gly Ile Arg Gly Leu Cys Asn Gly 65 ន

Lys Glu Leu Pro Ile Thr Gly Thr Asp Leu Gly Thr Asn Thr Ser Pro 100 22

Pro Pro Phe Leu 125 Gln His His Phe Glu Gln Asn Ser Ile Ile Fro Gly 115  $$\rm 120$ 8

Leu Val Ala Leu Phe Ile 140

Gly Ala Cys Ser Val Leu Leu Ala Leu 130

Phe

Ser Trp Arg Lys His 160 Ser 155 Ser Asn Leu Ser Leu Arg 150 쳝 Pro Glu His ' 145 35

Ser His Ser His Pro His Asn Thr Gln Ala Pro Gly Glu Ala 170 175

Cys Gly

Asn Val 185 뵱 Leu Gln Asp 18 2 Pro g Lya 6

INFORMATION FOR SEQ ID NO: 455: 3

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SEQUENCE DESCRIPTION: SEQ ID NO: 455: (A) LENGTH: 163 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear X;

S

Met Leu Gln Thr Ser Aen Tyr Ser Leu Val Leu Ser Leu Gln Phe Leu 1 15 15 Tyr Asp Leu Phe Val Asn Ser Phe Ser Glu Leu Leu Gln 20 30 Leu Ser 55

Lys Thr Pro Val 11e Gln Leu Val Leu Phe 11e 11e Gln Asp 11e Ala

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Val Leu Phe Asn Ile Ile Ile Ile Phe Leu Met Phe Phe Asn Thr Phe  $50 \,$ 

Val Phe Gln Ala Gly Leu Val Asn Leu Leu Phe His Lys Phe Lys Gly 65 70 75

Thr Ile Ile Leu Thr Ala Val Tyr Phe Ala Leu Ser Ile Ser Leu His  $95 \hspace{1.5cm} 95$ 

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Val Trp Val Met Asn Leu Arg Trp Lys Asn Ser Asn Ser 100 Phe Ile Trp

Thr Asp Gly Leu Gln Met Leu Phe Val Phe Gln Arg Leu Ala Ala Val \$125\$

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Leu Tyr Cys Tyr Phe Tyr Lys Arg Thr Ala Val Arg Leu Gly Asp Pro 130 140

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His Phe Tyr Gln Asp 145 Ser Leu Trp Leu Arg Lys Glu Phe Met Gln Val 150 155 160

Arg Arg Xaa

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Ξ SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 456:

(A) LENGTH: 46 umino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

X SEQUENCE DESCRIPTION: SEQ ID NO: 456:

35

Met Arg Ile Gln Val Phe Ile Leu Leu Gly Ala Gly Gly Thr Ser 1 15

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Ala His Thr Val Ala Xaa 65 70

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SEQUENCE CHARACTERISTICS:

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INFORMATION FOR SEQ ID NO: 459:

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Ser Ser Glu Leu Leu Glu Gln Leu Leu Ser Val Gln 50 55 60

Phe Val Trp Gln

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Leu Leu Pro Val Ile Leu Ala Val Phe Pro Asp Ser Ala Gln Ile Val 20 25 30

Pro Leu Leu Lys Pro Ile Pro Arg Pro Gln Pro Glu Val Ile Phe Pro 35 40 45

Met Leu Pro Pro Phe Ser Leu Val Tyr

Thr His Phe Leu Val Ala Ser 10 15

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SEQUENCE DESCRIPTION: SEQ ID NO: 458: (A) LENGTH: 70 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear 20

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INFORMATION FOR SEQ ID NO: 458:

(1) SEQUENCE CHARACTERISTICS:

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Phe Ala Val Ser Leu Ala Ala Lys Xaa 100 105

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Ser Leu Leu Ile Phe Ser Phe Gln Lys Thr Glu Ala Lys Leu Ile Val 85 90 95

His Pro Ala Ser Leu Leu Ile Val Phe Ala Thr Ser Ile Ser Glu Ser 65 70 75 80

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Lys Ser Ser

Ala Glu Ala Asp Gly Val

Leu Gln Pro Arg Arg 60

Ser Ser

Thr Asp Thr His Ile Cys Val Cys Val Cys Ile Tyr Leu Ser Ser Val
35 40 45

d C Ser

Ser Pro Thr Glu Thr Ser Glu Gln Ile Arg Glu Lys

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 105 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(2) INFORMATION FOR SEQ ID NO: 457:

Gln Phe Thr Lys Pro Pro Ser Leu Pro Leu Glu Pro Glu Pro Ala Val 20 25 30

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SEQUENCE DESCRIPTION: SEQ ID NO: 457:

Met Ser Tyr Leu Ala Phe Leu Tyr Met Thr Phe Asp Phe Cys Cys Leu 1 15

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Phe Ser Thr Val Tyr Ala Pro Ser Phe Lys Tyr Ile Cys Val His  $20 \hspace{1cm} 25 \hspace{1cm} 30$ 

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SEQUENCE DESCRIPTION: SEQ ID NO: 459:

(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 155 amino acids

Ser Glu Arg Met Asn Ala Leu Phe Val Gln Phe Ala Glu Val Phe Pro 35 40 45

Ala Leu Val Gly Leu Ala Lys Leu Ser Glu Glu Ile Ser Ala Pro Val 20 25 30

Met Ala Leu Leu Ser Val Leu Arg Val Leu Leu Gly Gly Phe Phe
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Ala Cys Ala Arg Ser Ile Phe Ile Phe Asp Glu Met Asp Lys Met His 175 Ala Gly Leu Ile Asp Ala Ile Lys Pro Phe Leu Asp Tyr Asp Leu 180 Val Asp Gly Val Ser Tyr Gln Lys Ala Met Phe Ile Phe Leu Ser Asn 195 Ala Gly Ala Glu Arg Ile Thr Asp Val Ala Leu Asp Phe Trp Arg Ser 210 Gly Lys Gln Arg Glu Asp lle Lys Leu Lys Asp lle Glu His Ala Leu  $235 \ \ \,$  235 Ser Val Ser Val Phe Asn Asn Lys Asn Ser Gly Phe Trp His Ser Ser 255 Leu lle Asp Arg Asn Leu lle Asp Tyr Phe Val Pro Phe Leu Pro Leu 260  $$260\,$ Glu Tyr Lys His Leu Lys Met Cys Ile Arg Val Glu Met Gln Ser Arg  $_{\rm 275}$ Gly Tyr Glu Ile Asp Glu Asp Ile Val Ser Arg Val Ala Glu Glu Met 290 Phe Phe Pro Lys Glu Glu Arg Val Phe Ser Asp Lys Gly Cys Lys 320 310 Met Ile Leu Thr Leu Leu Ser Val Val Ser Thr Met Ala Ser 1 5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 462: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 461 Lys Leu Asp Tyr Tyr Tyr Asp Asp 325 (A) LEWSTH: 14 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 5 emino acids (B) TYPE: emino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS INFORMATION FOR SEQ ID NO: 461: (2) INFORMATION FOR SEQ ID NO: 462: Met Leu Lys Cys Ile 1 5 Thr Val Phe Thr 일 당 3 S 9 2 20 25 23 8 32 <del>4</del> 45 S 8 Leu Leu Ala Gly Leu Leu Leu Val Mat Gly Pro 70 78 80 Leu Lys Val Phe Gly Tyr Gln Pro Asp Pro Leu Asn Tyr Gln Ile Ala 50 Pro Met Leu Gln Glu Ile Ser Asn Leu Phe Leu Ile Leu Leu Met Met 95 Gly Ala Ile Phe Thr Leu Ala Ala Leu Lys Glu Ser Leu Ser Thr Cys 100 lle Pro Ala Ile Val Cys Leu Gly Phe Leu Leu Leu Asn Val Gly 115 Gin Leu Leu Ala Gin Thr Lys Lys Val Val Arg Pro Thr Arg Lys Lys 130 Lys Leu Gly Arg Ala Val Leu Gly Leu Leu Leu Leu Ala Pro Ser 10 Leu Tyr Lys Asp Gln Leu Gln Leu Trp 11e Arg Gly Asn Val Ser 150 Val Val Gln Ala Val Glu Pro Ile Ser Leu Gly Leu Ala Leu Ala Gly 25 Val Leu Thr Gly Tyr Ile Tyr Pro Arg Leu Tyr Cys Leu Phe Ala Glu 15 45 Leu Asp Asp Asp Leu Phe Gly Gln His Leu Ala Lys Lys Ile Ile Leu 65 78 Asn Ala Val Phe Gly Phe lle Asn Asn Pro Lys Pro Lys Lys Pro Leu 95 Thr Leu Ser Leu His Gly Trp Thr Gly Thr Gly Lys Asn Phe Val Ser 110 Lys Ile Ile Ala Glu Asn Ile Tyr Glu Gly Gly Leu Asn Ser Asp Tyr 115 Val His Leu Phe Val Ala Thr Leu His Phe Pro His Ala Ser Asn Ile 130 Cys Cys Gly Gln Lys Arg Ser Leu Ser Arg Glu Ala Leu Gln Lys 50 -60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 460: Thr Leu Ser Thr Phe Lys Glu Ser Trp Lys Xaa 145 145 150 (A) LENGTH: 332 amino acids (B) TYPE: amino acid SEQUENCE CHARACTERISTICS: (2) INFORMATION FOR SEQ ID NO: 460 Val Gly Phe Leu Glu 65 ¥et 142 2 2 2 25 33 8 8 9 45 S 55

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Αla Ala Gly Tyr Gly Phe Ala Phe Arg Tyr Cys Pro Ser Gly Lys His Val 145 150 150 å Ser Gly Arg Ser Cys Thr Arg Pro His Cys Trp Pro Ser Leu Pro Ala 50 Ala Leu Thr Tyr Glu Glu Lys Pro Pro Tyr Ala Met Leu Arg Asn Asn 260 265 Gln Lys Phe Val Asp Lys Pro Gly Pro Phe Val Gly Pro Cys Gly His 225 230 230 ro Ę Ile 145 Ala Leu Glu Phe Leu His Glu Asn Glu Tyr Val His Gly Asn Val Thr 115 120 125 Ala Trp Gly Gly Ala Phe Ser Arg Pro Trp Met Ser Ala Gln Ser Met
90
95 8 4 Met Ser Lys Ser Ser His Ser Asn Trp Met Pro Arg Met Gly Ala Cys Ser 20 25 Met Lys Leu His Pro Pro Pro Pro Ser Pro Val Thr Gln Asp His Arg
1 15 Trp Ile Arg Pro Gln Ser Leu Gly Tyr Cys Met Leu Lys Trp Leu Tyr Gly Phe Leu 195 200 205 Tyr Val Glu Gly Ser Arg Ser Pro His Glu Gly Asp Leu Glu Phe  $165\,$ Cys Ala Glu Arg Ser Val Leu 100 Ser Trp Thr Asn Cys Leu Pro Xaa Xaa Glu Asp Ile Met Lys Gln Lys 210 225 Ser Met Asp Leu His Lys Gly Cys Gly Pro Ser Arg Arg Xaa Asp 180 185 190 Val Ser Val Ŷ. (i) SEQUENCE CHARACTERISTICS: Arg Thr Ser Ser Ser 35 SEQUENCE DESCRIPTION: SEQ ID NO: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 285 amino acids 뫈 Ser Glu Thr Leu Gln Lys Tyr Leu Lys Val Val Met 245 250 255 The Arg The A Gly Pro Gln Val Ala Cys Arg Leu Leu Asp 105 110 Asn Thr Gly Ser Trp Cys Tyr Pro 75 Pro Ser Leu Cys Lys Ser Thr 45

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Trp Gly Lys Arg Ile Ile Ser Glu His Cys Ser Ala Gln Ser Ser Xaa 65 70 80

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Ala Val Glu Arg Leu Leu Gly Val Arg Cys Thr Cys Pro Leu Ser 50 55

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Tyr Cys Leu Leu Val Ala Asn Gln Ser Ile Phe Phe Pro Cys Leu 35 40 45

Glu Ser Gln Val Pro Leu Ala Leu Ser Arg Val Phe Ser Thr Ser His 20 25 30

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SEQUENCE DESCRIPTION: SEQ ID NO: 464:

(B) TYPE: amino acid
(D) TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 80 amino acids

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(2) INFORMATION FOR SEQ ID NO: 464:

Leu Glu Ala Leu Leu Gln Asp Leu Arg Val Ser Pro 275

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INFORMATION FOR SEQ ID NO: 463:

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8 55 8 INFORMATION FOR SEQ ID NO: 466: £ SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 466: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 96 amino acids

8 Pro Leu Leu Val Lys Cys Arg Cly Arg Leu Lys Gly Val Asn Ile 35 40 45

5 Met His Thr Trp Tyr Asn Asp Arg Arg Gln Asn Cys His Cys Leu Leu 1 5 Phe Phe Leu Ile Tyr Leu Arg Lys Ile Tyr Gln Val Val Pro His Val 20 25 30

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 465: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENOTH: 47 amino acids

35 (2) INFORMATION FOR SEQ ID NO: 465:  $\Xi$ SEQUENCE CHARACTERISTICS:

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Met Glu Leu Val Leu Val Phe Leu Cys Ser Leu Leu Ala Pro Met Val 1 5 Leu Ala Ser Ala Ala Glu Lys Glu Het Asp Pro Phe His Tyr 25 Asp Tyr Gln Thr Leu Arg Ile Gly Gly Leu Val Phe Ala Val Val Leu 45 Phe Ser Val Gly Ile Leu Leu Ile Leu Ser Arg Arg Cys Lys Cys Ser 50 60 Asn Leu Ile Thr Ala Asn Ala Thr Glu Pro Gln Lys Ala Glu Asn Xaa 95 Phe Asn Gln Lys Pro Arg Ala Pro Gly Asp Glu Glu Ala Gln Val Glu 65 2 15

(2) INFORMATION FOR SEQ ID NO: 467: 25

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 467; (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 399 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear 8

Met Ala Ser Gly Ala Asp Ser Lys Gly Asp Asp Leu Ser Thr Ala Ile 1 5 10 Leu Lys Gln Lys Asn Arg Pro Asn Arg Leu Ile Val Asp Glu Ala Ile 20 Asn Glu Asp Asn Ser Val Val Ser Leu Ser Gln Pro Lys Met Asp Glu 15 Leu Gln Leu Phe Arg Gly Asp Thr Val Leu Leu Lys Gly Lys Lys Arg 50 60 35 <del>수</del>

Arg Glu Ala Val Cys Ile Val Leu Ser Asp Asp Thr Cys Ser Asp Glu 65 Lys Ile Arg Met Asn Arg Val Val Arg Asn Asn Leu Arg Val Arg Leu 95 45

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Gly Asp Val Ile Ser Ile Gln Pro Cys Pro Asp Val Lys Tyr Gly Lys 100 Arg Ile His Val Leu Pro Ile Asp Asp Thr Val Glu Gly Ile Thr Gly 116 115 55

Pro Ile Arg Lys Gly Asp Ile Phe Leu Val Arg Gly Gly Met Arg Ala 8

Asn Leu Phe Glu Val Tyr Leu Lys Pro Tyr Phe Leu Glu Ala Tyr Arg 130

160	Val	G]u	S.	Ter.	G1y 240	Æg	Pro	£.y3	Asp	Va.1 320	Lys	Ser	Asp
	11e 175	Arg	GL <sub>Y</sub>	Pro	Arg		Gly	Arg	Ile	Thr His Gly Glu Val 320	J35	Asn	
	Ş	Lys 190	Ile	Glu Leu Pro	Pro	Gly Lys Thr Leu Ile Ala 250 · 255	Asn 270	Lea	Phe	Gly	Gly	Pro 350	Glu Val
	Ser Pro Tyr	Gly Glu Pro Ile Lys 190	A8p 205		Pro	Fer.	. 11e	Asn 285	11e	H.s	Leu Met Asp Gly		Arg 365
	Pro	Pro	Asp	Val 220	Lys	Thr	Phe Leu	Ser	11e 300	큪	Met	Asn	Asp
155		Glu	Gly Tyr	Glu Met	Val 235	Lys	Phe	Glu	Ale	Lys 315	Zer Les	Ala Thr Asn Arg	Gly Arg Phe Asp
	Pro 170	G <u>L</u>			Gly Val 235	G1y 250	Phe	Ser	Pro	Glu Lys 315	330 330	Ala	Arg
	Glu Thr Asp	Glu 185	val	ile Lys	Ile	Pro Pro Gly Thr	Phe 265	GJu	Ala			Ala 345	Gly
	Thr	\$	G1u 200		Lys Ala	ΩŢ	Ala	G1y 280	Lys' Asn 295	Pro Lys Arg	Gln Leu Leu		Phe 360
	Glu	Ile His	Ser Leu Asn	Gln 215	Lys	Pro	Glu Thr Gly Ala				Gln	ile Val Met	
120	Val		3	Gln Leu Ala	Phe 230	Pro	뒾	Lys Leu Ala	Glu	Ala 310	Ser	116	Ala Leu Arg Arg
	Val 165	Val	Ser	<u> F</u>	Ala Leu	Gly 245	Glu		Ala	11e	Val 325	Val	Lea
	Lys	Ā 81 81	Glu		Ala	<u>ጟ</u>	Asn 260	Ser	Glu Glu	Ala	ile Val 325	His Val 340	
	Glu Phe Lys	Ala Pro Asp	Glu 195	Lys	Pro	lle Leu Leu Tyr	Ala	Met 275		Leu Asp Ala	Glu Arg Arg	Ala	Pro 355
	GF.	Pro	Glu	Arg 210	His	<u>F</u>	Ala val	116	Phe 290	<u>Leu</u>	Arg	Gln Arg	Asp
145	Val	Ala	Asp	2	Arg 225	Ile	Ala	Glu	Ala	Glu 305	Glu	Gln	11e
	<b>v</b> 1	•	10		15	20	i	25		30	35		40

(2) INFORMATION FOR SEQ ID NO: 468:

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lle Gly lle Pro Asp Ala Thr Gly Arg Leu Glu lle Leu Gln lle His 370 380

Thr Lys Asn Met Lys Leu Ala Asp Asp Val Asp Leu Glu Gln Xaa 385

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 468:

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2 INFORMATION FOR SEQ ID NO: 469:

£ SEQUENCE CHARACTERISTICS

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Œ. SEQUENCE DESCRIPTION: SEQ ID NO: 469: (A) LENGTH: 273 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Ala Ala Pro Lys Gly Ser Leu Trp Val Arg Thr Gln Leu Gly Leu 1 10

5

Ala Ser Ala Glu Ala Phe Asp Ser Val Leu Gly Asp 35Pro Pro Leu Leu Leu Thr Met Ala Leu Ala Gly Gly Ser Gly Thr  $20 \ \ 30$ o Thr Ala Ser

15

His Arg Ala Cys Gin Leu Thr Tyr Pro Leu His Thr Tyr Pro Lys Glu 50

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Gln Phe Val Asp Asp Gly Ile Asp Leu Asn Arg Thr Lys Leu Glu Cys 85 90 95 Glu Glu Leu Tyr Ala Cys Gln Arg Gly Cys Arg Leu Phe Ser Ile Cys 65 70 75 80

23

Thr Glu Lys Arg Leu Thr Lys Leu Leu Val His Ser Ser Leu Val Gly 50 55

Ser Ile Leu Ser Ala Leu Ser Ala Leu Val Gly Phe Ile Ile Leu Ser 65 70 75 80

Phe Ile Gly Pro Phe Phe Phe Ile Ile Ser Gly Ser Leu Ser Ile Ala 35  $40\,$ 

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Met Met Val Leu Ser Leu Gly Ile Ile Leu Ala Ser Ala Ser Phe Ser 1 15

SEQUENCE DESCRIPTION: SEQ ID NO: 470:

Pro Asn Phe Thr Gln Val Thr Ser Thr Leu Leu Asn Ser Ala Tyr Pro  $25\ 30$ 

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SEQUENCE CHARACTERISTICS

acids

(A) LENGTH: 192 amino (B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 470:

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Glu Ser Ala Cys Thr Glu Ala Tyr Ser Gln Ser Asp Glu Gln Tyr Ala 100 105 110

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Cys His Leu Gly Cys Gln Asn Gln Leu Pro Phe Ala Glu Leu Arg Gln 115 120 125

Glu Gln Leu Met Ser Leu Met Pro Lys 130 135 Met His Leu Leu Phe Pro Leu 140

Thr Leu Val Arg Ser Phe Trp Ser Asp Met Met Asp Ser Ala Gln Ser 145 150 150

Phe Ile Thr Ser Ser Trp Thr Phe Tyr Leu Gln Ala Asp Asp Gly Lys 165 170 175

Ile Val Ile Phe Xaa Ser Lys Pro Arg Asn Pro Arg Tyr Ala Pro His 180

Leu Glu Pro Gly Ala Leu Pro Asn Leu Xaa Xaa Xaa Ser Leu Ser Lys 200 205

Met

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Ser Xaa Xaa Ser Xaa Met Arg Asn Ser Gln Ala His Arg Asn Phe 210 220

Leu Glu Asp, Gly Glu Ser Asp Gly Phe Leu Arg Cys Leu Ser Leu Asn 225 230 230

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Ser Gly Trp Ile Leu Thr Thr Leu Val Leu Ser Val Met Val Leu 250 255

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Trp Ile Cys Cys Ala Thr Cys Cys Tyr Thr Leu Leu Asp Ala Val 260 265

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30 Val Lys Gln Ala Thr Leu Asn Pro Ala 85 Ser Leu Gln Cys Glu Leu Asp 90 95

Lys Asn Asn Ile Pro Thr Arg Ser Tyr Val Ser Tyr Phe Tyr His Asp 100 105

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Ser Leu Tyr Thr Thr Asp Cys Tyr Thr Ala Lys Ala Ser Leu Ala Gly
115 120 125

Xaa Leu Ser Leu Met Leu Ile Cys Thr Leu Leu Glu Phe Cys Leu Ala 130 140

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Val Leu Thr J 145 Ala val 150 Arg Trp Lys Gln Ala Tyr Ser Asp Phe Pro 155 160

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Gly Sex Val Leu Phe Leu Pro His Ser Tyr Ile Gly Asn Sex Gly Met
165 170 175 Ser 各 Met Thr His Asp Cys Gly Tyr Glu Glu Leu Leu Thr Ser 180 185

9 INFORMATION FOR SEQ ID NO: 471:

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 $\widehat{\Xi}$ SEQUENCE CHARACTERISTICS: (A) LENGTH: 234 amino acids
(B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 471:

Met Arg Lys Thr Arg Leu Trp Gly Leu Leu Trp Met Leu Phe Val Ser 1  $$\rm 10$ Glu Leu Arg Ala Ala Thr Lys Leu Thr Glu Glu Lys Tyr Glu Leu Lys 20

Glu Gly Gln Thr Leu Asp Val Lys Cys Asp Tyr Thr Leu Glu Lys Phe 35 2

Ala Ser Ser Gin Lys Ala Trp Gin Ile Ile Arg Asp Gly Glu Met Pro S0 60

Lys Thr Leu Ala Cys Thr Glu Arg Pro Ser Lys Asn Ser His Pro Val 65 78

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Gln Val Gly Arg Ile Ile Leu Glu Asp Tyr His Asp His Gly Leu Leu 95

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. Leu Tyr Gln 110 Arg Val Arg Met Val Asn Leu Gln Val Glu Asp Ser Gly 100

Cys Val Ile Tyr Gln Pro Pro Lys Glu Pro His Met Leu Phe Asp Arg 115 23

Ile Arg Leu Val Val Thr Lys Gly Phe Ser Gly Thr Pro Gly Ser Asn 130

Glu Asn Ser Thr Gln Asn Val Tyr Lys Ile Pro Pro Thr Thr Thr Lys 145 8

Ala Leu Cys Pro Leu Tyr Thr Ser Pro Arg Thr Val Thr Gln Ala Pro 175 Pro Lys Ser Thr Ala Asp Val Ser Thr Pro Asp Ser Glu Ile Asn Leu 180

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Thr Asn Val Thr Asp ile Ile Arg Val Pro Val Phe Asn Ile Val Ile 200 <del>8</del>

Thr Leu Arg Ser Phe Val Pro 230 Phe Ala Val

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(2) INFORMATION FOR SEQ ID NO: 472:

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SEQUENCE CHARACTERISTICS:
(A) LENGTH: 105 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Met Leu His Ile Leu Pro Leu Lys Ser Tyr Asp Phe Pro His Phe Ser 10 8

SEQUENCE DESCRIPTION: SEQ ID NO: 472.

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Leu Met Gly Arg Tyr Arg Cys Ala Ser Leu Leu Phe Cys Phe Leu Leu .  $_{\rm 25}$ 

Leu Phe Phe Phe Cys Ser Val Leu Trp Thr Phe Ser Asp Met His S

Arg Ser Gly Glu Asp Gly Pro Trp Thr Pro Cys Val His His Leu Ala 50 60

Phe Ser Pro Val Leu Phe Ile Glu Asn Pro Arg His Tyr Ala Asn Ala 85 Ala Ser Leu Ile Ser Tyr Gly Gln Pro Gly Phe Ile Cys Ile Ser Leu 65 2

Thr Val Thr Thr Leu Gly Asp Trp Xea 100

2

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(2) INFORMATION FOR SEQ ID NO: 473:

(A) LENGTH: 32 amino acids (B) TYPE: amino acid (i) SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 473:

Met Val Phe Leu Lys Tyr Arg Phe Leu Phe Phe Leu Val Phe Leu Ala  $1 \ \ \, 1$ Asn Cys Ila Tyr Ser Leu His Tyr Lys Pro Ser Leu Met Tyr Pro Lys 20 25 2

<del>8</del>

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(2) INFORMATION FOR SEQ ID NO: 474:

(A) LENGTH: 571 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 474:

Met Ala Leu Ser Arg Gly Leu Pro Arg Glu Leu Ala Glu Ala 15 Gly Gly Arg Val Leu Val Val Gly Ala Gly Gly Ile Gly Cys Glu Leu 20 30 ೱ

Leu Lys Asn Leu Val Leu Thr Gly Phe Ser His Ile Asp Leu Ile Asp 40 55

Leu Asp Thr Ile Asp Val Ser Asn Leu Asn Arg Gln Phe Leu Phe Gln 50 55 8

Lys Lys His Val Gly Arg Ser Lys Ala Gln Val Ala Lys Glu Ser Val 65 70 75 80

Leu Gln Phe Tyr Pro Lys Ala Asn Ile Val Ala Tyr His Asp 85 Ser Ile 95

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Met Asn Pro Asp Tyr Asn Val Glu Phe Phe Arg Gln Phe Ile Leu Val 105

Met Asn Ala Leu Asp Asn Arg Ala Ala Arg Asn His Val Asn Arg Met 115 120 125

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Cys Leu Ala Ala Asp Val 130 Pro Leu Ile Glu Ser Gly Thr Ala Gly Tyr 135

2 Leu Gly Gln Val Thr Thr Ile Lys Lys Gly Val Thr Glu Cys Tyr Glu 145 150 155 160

CYS His Pro Lys Pro Thr Gln Arg Thr Phe Pro Gly Cys Thr Ile Arg 165 170

20

Asn Thr Pro

Ser Glu Pro Ile His Cys Ile Val Trp Ala Lys Tyr Leu 180 190

25

Phe Asn Gln Leu Phe Gly Glu Glu Asp Ala Asp Gln Glu Val Ser Pro 195 200 205

Asp Arg Ala Asp Pro Glu Ala Ala Trp Glu Pro Thr Glu Ala Glu Ala 210 215

30

Leu Ile Asn Ile Leu 530

His

Ser Glu Asp Leu Gly Lys Asp Val Glu Phe 535

Glu Val Val Gly Asp Ala Pro Glu Lys Val Gly Xaa Lys Gln Ala 545 550 555

560 51u

25

Asn Gly Ser Arg Leu Gln Ala Asp Asp Phe Leu Gln Asp Tyr Thr Leu 515 526 525

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The Glu

Ala Asn Asn His Lys Lys Leu Ser Glu Phe Gly Ile Arg 500 505 510

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Gln Asp Lys Ile Val Lys Glu Lys Phe Ala Met Val Ala Pro Asp Val 465 470 475

Glu Val Thr Val Arg Leu Asn Val His Lys Val Thr Val Leu Thr Leu 450 460

Ala Leu Asp Pro Pro Asn Pro Asn Cys Tyr Val Cys Ala Ser Lys 435 440 445

Pro

Phe Leu Asn Lys Gln Pro Asn Pro Arg Lys Lys Leu Leu Val Pro Cys 420 425

Glu Gly Leu Lys Ile Leu Ser Gly Lys Ile Asp Gln Cys Arg Thr Ile 415

Pro Ala Ile Ala Thr Thr Asn Ala Val Ile Ala Gly Leu Ile Val Leu 385 390 390

Gln Ile Glu Asp Gly Lys Gly Thr Ile Leu Ile Ser Ser Glu Glu Gly 495

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Arg Ala Arg Ala Ser Asn Glu Asp Gly Asp Ile Lys Arg Ile Ser 225 230 235 246 246

Lys Glu Trp Ala Lys Ser Thr Gly Tyr Asp Pro Val Lys Leu Phe Thr  $245\ 250\$ 

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Leu Phe Lys Asp Asp Ile Arg Tyr Leu Leu Thr Met Asp Lys Leu 260 265 270

35 Asp Ala Ala Lys Ser Ile Thr Asn Gly Gln Xaa 565 570

(i) SEQUENCE CHARACTERISTICS

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INFORMATION FOR SEQ ID NO: 475:

(A) LENGTH: 312 amino acids
(B) TYPE: amino acid

Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 475: (D) TOPOLOGY: linear

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Met Oln Val Val Thr Cys Leu Thr Arg Asp Ser Tyr Leu Thr His Cys
1 10 15

Phe Leu Gln His Leu Met Val Val Leu Ser Ser Leu Glu Arg Thr Pro 20 25 30

8

Ser Pro Glu Pro Val Asp Lys Asp 35 Phe Tyr Ser Glu Phe Gly Asn Lys 45

Thr Thr Gly Lys Met Glu Asn Tyr Glu Leu Ile His Ser Ser Arg Val 50 60

Lys Phe Thr Tyr Pro Ser Glu Glu Glu Ile Gly Asp Leu Thr Phe Thr 65 70 75

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Asn Met Lys Ser Arg Phe Asp Ile Lys Ser Met Ala Gly Asn Ile Ile 370 , 375 , 380

Asp Phe Val Thr Ser Ala Ala Asn Leu Arg Met His Ile Phe Ser Met 355 360 365

Gly Asp Gly Ala Glu Leu Ile Trp Asp Lys Asp 340 345

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Pro Ser Ala Met 350

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Phe

Ser Lys Ser Ile Glu Thr Leu 325

Arg Val His Leu Ala Glu Lys 330 335

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Ser Gin Gly Glu Glu Thr Asn Ala Ser Asp Gin Gin Asn Glu Pro Gin 290 100

Leu Gly Leu 305

Lys Asp Gln Gln Val Leu Asp Val Lys Ser Tyr Ala Arg 310 315

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Lys Arg Lys Pro Pro Val Pro Leu Asp Trp Ala Glu Val Gln 275 280 285

52

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Val Ala Gin Lys Het Ala Giu Pro Giu Lys Ala Pro Ala Leu Ser Ile 95 5 Leu Leu Tyr Val Gin Ala Phe Gin Val Gly Het Pro Pro Pro Gly Cys

Leu Leu Tyr Val Gln Ala Phe Gln Val Gly Met Pro Pro Gly Cys 100 Cys Arg Gly Pro Leu Arg Pro Lys Thr Leu Leu Leu Thr Ser Ser Glu 115

9

Ile Phe Leu Leu Asp Glu Asp Cys Val His Tyr Pro Leu Pro Glu Phe 130

Ala Lys Glu Pro Pro Gln Arg Asp Arg Tyr Arg Leu Asp Asp Gly Arg 145 160

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145 155 150 155 160 Arg Val Arg Val Arg Val Arg Asp Leu Asp Arg Val Leu Met Gly Tyr Gln Thr Tyr Pro 175 170 175

20 Gln Pro Ser Pro Ser Ser Met Thr Cys Lys Val Met Thr Ser Trp 180 190

Ala Val Ser Pro Trp Thr Thr Leu Gly Arg Cys Gln Val Ala Arg Leu 195

25 205 205 Clu Pro Ala Arg Ala Val Lys Ser Ser Gly Arg Cys Leu Ser Pro Val 210 210

Leu Arg Ala Glu Arg Ser Ser Arg Cys Trp Leu Ala Ser Gly Arg 230 225 235 230

Gin Ala Phe Val Phe Ser Lys Asn Val Leu Ser Ser Leu Trp Tyr Leu 275

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Asn Leu Thr Val Leu Ala Glu Asn Val Asn Met Cys Val Cys Cys Val 290

Asn Ser Phe Ser Cys Trp Glu Xaa 310

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(2) INFORMATION FOR SEQ ID NO: 476:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 329 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 476:

Met Ala Gln His His Leu Trp Ile Leu Leu Cys Leu Gln Thr Trp

1 5 10 15

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60 Pro Glu Ala Ala Gly Lys Asp Ser Glu Ile Phe Thr Val Asn Gly Ile

The Gin Ala Asp Pro Tyr The The Lys Arg Tyr Asn Leu Gin Ile
115
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17/r Arg Arg Leu Gly Lys Pro Lys Ile The Gin Ser Leu Met Ala Ser
130
130

130 135 140

Val Asn Ser Thr Cys Asn Val Thr Leu Thr Cys Ser Val Glu Lys Glu

150 145 155 160

Glu Lys Asn Val Thr Tyr Asn Trp Ser Pro Leu Gly Glu Gly Asn

165 175

30 Val Leu Gln Ile Phe Gln Thr Pro Glu Asp Gln Glu Leu Thr Tyr Thr 185
180
180
180
Cys Thr Ala Gln Asn Pro Val Ser Asn Asn Ser Asp Ser Ile Ser Ala
195
205
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Arg Gln Leu Cys Ala Asp Ile Ala Met Gly Phe Arg Thr His His Thr
210
215

Gly Leu Leu Ser Val Leu Ala Met Phe Phe Leu Leu Val Leu Ile Leu 40 215 240 240 245 Ser Ser Val Phe Leu Phe Arg Leu Phe Lys Arg Arg Gln Asp Ala Ala 245 245

45 Ser Lys Lys Thr Ile Tyr Thr Tyr Ile Net Ala Ser Arg Asn Thr Gln 260 270 Pro Ala Glu Ser Arg Ile Tyr Asp Glu Ile Leu Gln Ser Lys Val Leu 275 286 286 286 286 Pro Ser Lys Glu Glu Pro Val Asn Thr Val Tyr Ser Glu Val Gln Phe 290 390

Ala Asp Lys Met Gly Lys Ala Ser Thr Gln Asp Ser Lys Pro Gly 55 305

Thr Ser Ser Tyr Glu Ile Val Ile Xaa

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Gin Ala Lys Asp Thr Phe Pro Asn Val Thr Ile Leu Ser Thr Leu Glu 50 60 Asp Met His His Ile Glu Glu Ser Phe Gln Glu Ile Lys Arg Ala Ile \$45\$Leu Cys Ser Val Asp Asn His Gly Leu Arg Arg Cys Leu Ile Ser Thr  $20 \ \ 30$ Met Lys Leu Gin Cys Val Ser Leu Trp Leu Leu Gly Thr Ile Leu Ile 1 15 Cys Arg Gln Glu Ala Thr Asn Ala Thr Arg Val Ile His Asp Asn Tyr 130 140 Tyr Met Gln Lys Thr Leu Arg Gln Cys Gln Glu Gln Arg Gln Cys His 115 120 125 Leu Leu Ala Phe Tyr Val Asp Arg Val Phe Lys Asp His Gln Glu Pro 85 90 Asp Val Phe Leu Ala Trp Ile Asn Lys Asn His Glu Val Met Ser Ser 175 Asn Pro Lys Ile Leu Arg Lys Ile Ser Ser Ile Ala Asn Ser Phe Leu 100 105 Thr Leu Gin Ile Ile Lys Pro Leu Asp Val Cys Cys Val Thr Lys Asn 65 70 75 80 Asp Gin Leu Giu Val His Ala Ala Ala Ile Lys Ser Leu Gly Giu Leu 145 150 150 2 Ala Xaa Asp Thr Ala Ile Arg Val Ala Leu Ala Val Ala Val Leu Lys Thr Val 1 15 The Leu Gly Leu Leu Cys Cly Gly Gly Gly Gly Gly Gly Lys
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30 INFORMATION FOR SEQ ID NO: 478: (i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 52 amino acids (x1)' SEQUENCE DESCRIPTION: SEQ ID NO: 478: (B) TYPE: amino acid

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35 30 25 20 15 5 Arg Asp Val Tyr 50 Met Gln Lys Lys Asn Ser Leu Phe Phe Phe Phe Ala Phe Tyr Tyr Glu 10 Ala Ile Asn Lys Leu Asn Tyr Leu His Trp Thr His Phe Gln  $_{50}$ Lys Glu Tyr Phe Leu Pro 35 Asn Lys Thr Asn Ala Pro Gly Glu Gly Ser Met Ile Thr Arg Asn Ile 20 25 2 (2) INFORMATION FOR SEQ ID NO: 480: INFORMATION FOR SEQ ID NO: 479: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 479: (i) SEQUENCE CHARACTERISTICS (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 62 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 27 amino acids Phe Leu Phe Cys Cys Val Glu Ala Ser Ile 40 45

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SEQUENCE DESCRIPTION: SEQ ID NO: 477: (A) LENGTH: 178 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 477:

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(1) SEQUENCE CHARACTERISTICS (B) TYPE: amino acid (A) LENGTH: 339 amino acids

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INFORMATION FOR SEQ ID NO: 481:

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Met Ser Gly Pro Asp Val Glu Thr Pro Ser Ala Ile Gln Ile Cys Arg 1 10 15 Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 481: (D) TOPOLOGY: linear

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8 Ile Met Arg Pro Asp Asp Ala Asn Val Ala Gly Asn Val His Gly Gly
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Val Ala Gly Arg Gln Ala Val Thr Ser Asp Gln Gln Ser Val Gly Arg
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(ž SEQUENCE DESCRIPTION: SEQ ID NO: 480: (D) TOPOLOGY: linear

Met Pro Gly Leu Ser Leu Ile Leu Thr Val Thr Leu Leu Ala Val Ser 1 15

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Asp Ser Ala Ala Thr Cys Ile Val Ala Lys Gly
20
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Thr Ile Leu Lys Met Ile Glu Glu Ala Gly Ala Ile Ile Ser Thr Arg 40 Gln Val Ann Val Met Ser Glu Asn Ile Leu Thr Gly Ala Lys Lys Leu 100 Phe Leu Ser Pro Met Cys Ile Gly Glu Val Ala 70 75 80 Lys His Ser Val Glu Val His Cys Asn Ser Gln Asn Gly Glu Arg Cys Val Ala Ala Leu Ala Arg Lys Val Leu Glu Val Pro Pro Val Val Tyr Ser Arg Xea Glu Gln 130 Glu Thr Lys Trp Arg Asn Gly Asp Ile Val Gln Pro Val Leu Asn Pro 175 Val Val Asp Ser Ser Gin Lys Arg Tyr Arg Ala Ala Ser Ala Phe Phe 275 Leu Val Pro Glu Thr Glu Asp Glu Lys Lys Arg Phe Glu Glu Gly Lys 320 315 Gly Arg Tyr Leu Gln Met Lys Ala Lys Xaa Gln Gly His Ala Xaa Xaa 335 Thr Asn Lys Ala Thr Leu Trp Tyr Val Pro Leu Ser Leu Lys Asn Val 115 Glu Glu Glu Gly Arg Lys Arg Tyr Glu Ala Gln Lys Leu Glu Arg Met 165 Glu Pro Asn Thr Val Ser Tyr Ser Gln Ser Ser Leu Ile His Leu Val 180 Gly Pro Ser Asp Cys Thr Leu His Gly Phe Val His Gly Gly Val Thr 195 Lys Leu Met Asp Glu Val Ala Gly Ile Val Ala Ala Arg His Cys 210 Lys Thr Asn Ile Val Thr Ala Ser Val Asp Ala Ile Asn Phe His Asp 225 Lys Ile Arg Lys Gly Cys Val Ile Thr Ile Ser Gly Arg Met Thr Phe 255 Thr Ser Asn Lys Ser Met Glu Ile Glu Val Leu Val Asp Ala Asp Pro 260 Thr Tyr Val Ser Leu Ser Gln Glu Gly Arg Ser Leu Pro Val Pro Gln 290 His Val Ser Ala Glu Ile Thr Tyr Thr Ser 85 90 Val Glu Arg Thr Asp Agp Met 65 S 2 2 2 8 35 23 6 45 S 55

(2) INFORMATION FOR SEQ ID NO: 482:

(A) LENGTH: 32 amino acids (i) SEQUENCE CHARACTERISTICS:

TYPE: amino acid

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Met Leu Asn Ser Asn Ile Asn Asp Leu Leu Met Val Thr Tyr Leu Ala 1  $10\,$ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 482:

Asn Leu Thr Gin Ser Gln Ile Ala Leu Asn Glu Lys Leu Val Asn Leu 20 15

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(2) INFORMATION FOR SEQ ID NO: 483:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 48 amino acids

22

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 483:

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Met Arg Glu Thr Ser Ile Arg Val Leu Leu Met Leu Pro Ala Leu Glu 1 1 15 15

Cys Phe Lys Thr Ile Thr Cys Trp Pro Thr Ser Leu Thr Gln Arg Xaa  $_{\rm AS}$ Ser Thr Ser Gly Leu Ser Ala Phe Met Gly Leu Gly Thr Arg Ile Gly 25

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(2) INFORMATION FOR SEQ ID NO: 484: 45 (i) SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 484:

S

Met Tyr Met Tyr Ser Leu Asn Val Phe Leu Ser Phe Ile Phe Leu Ala 1 5

Leu Val Phe Lys Cys Val His Val Cys Gln Gly Ala Asn Ala Phe Leu 20

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Phe Leu Lys Leu Val Phe 35

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Gln Pro Xaa

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INFORMATION FOR SEQ ID NO: 485:

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30 20 2 5 2. 6 325 S 50 Leu Val Phe Ala Trp Glu Phe Phe Ser Glu Asp Thr Pro 50 55 60 Gln Glu Cys Thr Glu Lys Phe Ala Lys Leu Leu Val Gln Leu Ile Ser 35 40 Val Cys Glu Met Phe Leu Phe Phe Leu Met Thr Gln Lys Leu Ile Trp \$20\$Met Gly Leu Arg Leu Ile Cys Leu Glu Leu Thr Met Val Lys Ala Leu 1 15 Thr Lys Asn Gin Trp Leu Leu Thr Pro Ser Arg Glu Tyr Ala Thr Lys  $35 \ \ 40 \ \ \ 45$ Met Leu Ala Ala Arg Leu Val Cys Leu Arg Thx Leu Pro Ser Arg Val 1 10 15 The Arg Ile Gly Ile Arg Arg Gly Arg The Gly Gln Glu Leu Lys Glu  $50 \ \mbox{50}$ Ala Val Ile Trp Pro Gln Tyr Val Lys Asp Arg Ile His Ser Thr Tyr 115 120 125 Gly Arg Trp Phe Val Ala Gly Gly Ala Ala Val Gly Leu Gly Ala Leu 85 90 95 Ala Ala Leu Glu Pro Ser Met Glu Lys Ile 65 70 Phe His Pro Ala Phe Thr Lys Ala Ser Pro Val Val Lys Asn Ser Ile 20 30 (2) INFORMATION FOR SEQ ID NO: 486: Met Tyr Leu Ala Gly Ser Ile Gly Leu Thr Ala Leu Ser Ala Ile Ala 130  $_{\dagger}$  135 140 Tyr Tyr Gly Leu Gly Leu Ξ Ξ ž (X) SEQUENCE CHARACTERISTICS SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear SEQUENCE DESCRIPTION: SEQ ID NO: 486: SEQUENCE DESCRIPTION: SEQ ID NO: 485: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 346 amino acids Ser Asn Glu Ile Gly Ala Ile Glu Lys 105 Phe Lys Ile Asp Gln Met 75

છ 20 5 6 25 5 35 2 8 55 50 Ile Val Gly Gly Leu Ser Thr Val Ala Met Cys Ala Pro Ser Glu Lys 225 230 230 235 Ile Leu Gly Gly Pro Leu Leu Ile Arg Ala Ala Trp Tyr Thr Ala Gly 210 215 220 Trp Leu Leu His Ser Gly Val Met Gly Ala Val Val Ala Pro Leu Thr 195 200 205 Val Arg Ser Ile Pro Tyr Asp Gln Ser Pro Gly Pro Lys His Leu Ala 180 185 Val Thr Ile Gly Val Thr Phe Ala Ala Met Val Gly Ala Gly Mat Leu 165 170 Ser Pro Met Tyr Gly Val Gln Lys Tyr Asp Pro Ile Asn Ser Met Leu 305 310 315 Ala Thr Leu Tyr Ser Val Ala Met Tyr Gly Gly Leu Val Leu Phe Ser 275 280 Ile Ser Arg Thr Pro Val Leu Met Asn Phe Met Met Arg Gly Ser Trp 145 150 Met Glu Glu Val Leu Leu Gly Leu Lys Asp Arg Glu Gly Tyr Thr 1 5 10 Leu Ala Thr Gly Gly Asn Arg Lys Lys Xaa 340 345 Ser Ile Tyr Met Asp Thr Leu Asn Ile Phe Met Arg Val Ala Thr Met 325 330 335 æc Val Ser Phe Leu Asn Met Gly Ala Pro Leu Gly Val Gly Leu Gly Leu Val Phe 245 250 255 Arg Arg Lys Ser Leu Leu Thr Arg Lys Val Ile Cys Lys Ser Asp Ala 50 55 60 Ile Glu Leu Ala Leu Arg Gly Arg Leu Gln Leu Glu Ala Cys Gly Met
35 40 45 (2) INFORMATION FOR SEQ ID NO: 487: Phe Leu 290 Phe Trp Ser Leu Gly Ser Met 260 Ě (i) SEQUENCE CHARACTERISTICS 5 Asn Asp Cys Ile Ser Ser Gly Leu Arg Gly Cys Met Leu 20 25 30 SEQUENCE DESCRIPTION: SEQ ID NO: 487: (A) LENGTH: 237 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear Tyr Asp Thr Gln Lys Val Ile Lys Arg Ala Glu Val  $295\,$ Phe Leu Pro Pro Thr Thr Val Ala Gly 265 270

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Asp Ile Ser Glu Ser Ile Lys Glu Leu Gln Phe Tyr Arg Asn Asn Ile 170 Glu Asn Glu Lys Thr Val Ser 195 2 2 2 25 8 35 <del>6</del> S 55 8 45 Ser Gly 95 Pro Thr Gly Asp Val Leu Leu Asp Glu Ala Leu Lys His Val Lys Glu 65 75 80 Pro Leu Lys Leu His Tyr Gln Leu Arg Asn Val Arg 105 Glu Arg Leu Ala Lys Asn Leu Val Glu Lys Gly Val Leu Thr Thr Glu 115 Leu Phe Asp Met Thr Thr His Pro Leu Thr Asn 115 15 E Asp Lys Trp Val Asn Asp Pro His Arg Met Asp Arg Arg Leu Leu Ala 175 Leu Ile Tyr Leu Ala His Ala Ser Asp Val Leu Glu Asn Ala Phe Ala 180 Pro Leu Leu Asp Glu Gln Tyr Asp Leu Ala Thr Lys Arg Val Arg Gln 200 Leu Asp Leu Asp Pro Glu Val Glu Cys Leu Lys Ala Asn Thr Asn 210 Ile Glu Lys Asp Gln Ile Ile Glu Met Ala Cys Leu Ile Thr Asp Ser 20 Asp Leu Asn Ile Leu Ala Glu Gly Pro Asn Leu Ile Ile Lys Gln Pro Ser Met Ser Asp Trp Cys Lys Glu His His Gly 55 Lys Ala Val Lys Glu Ser Thr Ile Thr Leu Gln 70 75 80 Asn Asn Ile Lys Gln Arg Leu Ile Lys Lys Val Gln Glu Ala Val 145 Š Met Ala Glu Arg Met Val Trp Val Asp Leu Glu Met Thr Gly 1  $$\rm 10$ ž Glu Val Leu Trp Ala Val Val Ala Ala Phe Thr Lys Xaa 225 (A) LENGTH: 200 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 488: Thr Gln Pro Pro Glu Thr Val Gln Asn Trp Ile Glu 85 90 (2) INFORMATION FOR SEQ ID NO: 488: Lys Ser Gly Leu Thr 65 Lys Gln Asn Phe Leu 130 Asp Glu Leu Leu Asp 50 Glu Thr Trp Asn 100 圣 8 2 2 8 6 45 ල 25 33 S 55

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Gly Leu Cys Pro Leu Ala Gly Asn Ser Val His Glu Asp Lys Lys Phe  $100\,$ Leu Asp Lys Tyr Met Pro Gln Phe Met Lys His Leu His Tyr Arg Ile 115

ile Asp Val Ser Thr Val Lys Glu Leu Cys Arg Arg Trp Tyr Pro Gly 130 Glu Tyr Glu Phe Ala Pro Lys Lys Ala Ala Ser His Arg Ala Leu Asp 145 Phe Lys Lys Lys Ile Asp Glu Lys Lys Arg Lys Ile Ile Glu Asn Gly 180

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(2) INFORMATION FOR SEQ ID NO: 489:

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 351 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Ala Thr Thr Ala Ala Pro Ala Gly Gly Ala Arg Aen Gly Ala Gly 1 5 10 15

SEQUENCE DESCRIPTION: SEQ ID NO: 489

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Val Ile Asp Met Glu Asn Met Asp Asp Thr Ser Gly Ser Ser Phe Glu 15 Asp Net Gly Glu Leu-His Gln Arg Leu Arg Glu Glu Glu Val Asp Ala 50 Asp Ala Ala Asp Ala Ala Ala Ala Glu Glu Glu Asp Gly Glu Phe Leu 65 Gly Met Lys Gly Phe Lys Gly Gln Leu Ser Arg Gln Val Ala Asp Gln 89 90 95 Met Trp Gin Ala Gly Lys Arg Gin Ala Ser Arg Ala Phe Ser Leu Tyr 100

Val Arg Thr Gly Leu Leu Glu Ser Met Ile Pro Ile Lys Met Val Asn 130 130 Ala Asn 11e Asp 11e Leu Arg Pro Tyr Phe Asp Val Glu Pro Ala Gln 115

Gln Ala Glu Tyr Glu Phe Leu Ser Phe Val Arg Gln Gln Thr Pro Pro

Phe Pro Gin Lys Ile Ala Gly Glu Leu Tyr Gly Pro Leu Met Leu Val 145 150 150

Phe

Thr Leu Val Ala Ile Leu Leu His Gly Met Lys Thr Ser Asp Thr 165 170 175

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55 50 8 Glu Ser Gly Thr Ser Gly Gly Gly Gly Ser Thr Glu Glu Ala Phe Met . 20  $$25\,$ Met Arg Gly Ser Arg Gly Gly Trp Ala Gly Glu Met Ala Ala Ser Gly . 1  $$\rm 10$ 컱 Thr Ser Lys Asn Glip Ile Glu Arg Leu Thr Arg Pro Gly Ser Ser Tyr 50 55 Phe Tyr Ser Glu Val Lys Gln Ile Glu Lys Arg Asp Ser Val Leu 35 40 45

(2) INFORMATION FOR SEQ ID NO: 490: Ë

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Leu Leu Tyr Leu His Phe Ala Tyr His Lys Val Val Glu Gly Ile Leu 290 300

Asp Thr Leu Glu Gly Pro Asn Ile Pro Pro Ile Gln Arg Val Pro Arg 305 310

Gln Arg Leu Leu Ceu Cys Gly Thr Leu Ala Ala Leu His Met Leu Phe 275 280 285

Thi Leu Arg Met Val Ala Val Leu Val Ser Arg Thr Val Gly Pro Thr  $260\ 265$ 

Leu His Ala Leu Phe Tyr Leu Phe Trp Leu Leu Val Gly Gly Leu Ser 245 250

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Asp Ile Pro Ala Met Leu Pro Ala Ala 325

Arg Leu Pro Thr Thr Val Leu 330

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Ala Thr Ala Lys Ala Val Ala Val Thr Leu Gln Ser His Xaa 340 345

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Gly Leu Phe Gly His Cys Ile Val Leu Phe Ile Thr Tyr Asn Ile His 225 230 230 235

Cys Asn Ala Gln Ile Thr Met Leu Gln Met Leu Ala Leu Leu Gly Tyr 210 220

Gly Tyr Trp Leu Gly Val Ser Ser Phe Ile Tyr Phe Leu Ala Tyr Leu 200 205

Ile Ile Arg Glu Gly Thr Leu Met Gly Thr Ala Ile Gly Thr Cys Phe 180 180

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X i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 265 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear SEQUENCE DESCRIPTION: SEQ ID NO: 490:

ᅜ 5 6 ဗ 25 20 ઝ Glu Ser Arg Asp Gly Arg Val Asp Ser Trp 230 Val Glu Glu Asp Asp Pro Glu Leu 165 Val Lys Glu Arg Lys Lys Gln Leu Lys Lys Glu Gly Lys Pro Thr Ile 145 150 150 Arg Ala Leu Asp Val Ile Gln Ala Gly Lys Glu Tyr Val Glu His Thr  $130\,$ Ala Val Asp Lys Ala Tyr Lys Leu Leu Leu Asp Gln Glu Gln Lys Lys 115 Pro Asp Lys Asn Gln Asp Asp Ala 100 Asp Glu Glu Ile Lys Lys Arg Phe Arg Gln Leu Ser Ile Leu Val His 85 90 95 Phe Asn Leu Asn Pro Phe Glu Val Leu Gln Ile Asp Pro Glu Val Thr 65 70 75 80 Lys Val Lys Met Glu Gln Arg Glu Xaa 260 265 Thr Lys Gly Lys Lys Glu Lys Lys Asn Arg Thr Phe Leu Arg Pro Pro 255 Ala Gin Glu Lys Ala Lys Arg Glu Arg Glu Trp Gln Lys Asn Phe Glu 210 220 Lys Met Lys Glu Met His Glu Arg Lys Arg Gln Arg Glu Glu Glu Ile Glu 195 200 Leu Phe Ala Glu Leu Glu Ile Lys Arg Lys Glu Arg Glu 180 185 190 Phe Lys Gln Ala Val Tyr Lys Gln 175 105 105 Arg Ala Gln Lys Ala Phe Glu Arg Asn Phe Gln Ala Asn 235 240

2 INFORMATION FOR SEQ ID NO: 491:

Ξ Ě SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 491: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 25 amino acids

25

Asp Ser Met Pro Thr Cys Pro Leu Xaa Ala Ser Leu Glu Cys Gly Pro 1 15

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55 Leu Leu Pro Val Arg Leu Cys Cys Leu 20

(2) INFORMATION FOR SEQ ID NO: 492:

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SEQUENCE DESCRIPTION: SEQ ID NO: (A) LENGTH: 159 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 3

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Asn Glu Tyr Arg Val Pro Glu Leu Asn Val Gln Asn Gly Val Leu 5

Ser Phe Leu Phe Glu Tyr Ile Gly Glu Met Gly Lys Asp  $20 \ \ \, 25$ Ş Ser Lys 2

Pro Leu Leu Glu Asp Ala Leu Met Asp Arg 40 Tyr Ile Tyr Ala Val Thr

Leu Val His Arg Gln Thr Ala Ser Ala Val Val Gln His Met Ser 50 60 Asp 2

Gly Val Tyr Gly Phe Gly Cys Glu Asp Ser Leu Asn His Leu Leu 70 75 58 63

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22

Asn Tyr Val Trp Pro Asn Val Phe Glu Thr Ser Pro His Val Ile Gln

Ala Val Met Gly Ala Leu Glu Gly Leu Arg Val Ala Ile Gly Pro Cys 100 Met Leu Gln Tyr Cys Leu Gln Gly Leu Phe His Pro Ala Arg Lys 115 Arg

Ser Arg Asp Val.Tyr Trp Lys lie Tyr Asn Ser lie Tyr lie Gly 130 ۷aJ 3

Leu Ile Ala His Tyr Pro Arg Ile Tyr Gln Arg Xaa 150 Gln Asp Ala I 145 35

INFORMATION FOR SEQ ID NO: 493: 3 6

(A) LENGTH: 279 amino acids TYPE: amino acid SEQUENCE DESCRIPTION: SEQ ID NO: 493:  $\widehat{\mathbf{x}}$ 

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Met Ile Ser Asp Asn Ser Ala Glu Asn Ile Ala Leu Val Thr Ser Met 1 15

Gly Leu Leu Gln Ala Gly Ala Arg Leu Cys Pro Thr Val Gln 20 35 Tyr Asp 20

Ala Lya Glu Gly Lya Ile Glu Ile Phe Arg His Ile Leu Gln Arg Glu 50 60 Leu Glu Asp Ile Arg Asn Leu Gln Asp Leu Thr Pro Leu Lys Leu Ala 15 55

Ser Gly Leu Ser His Leu Ser Arg Lys Phe Thr Glu Trp Cys Tyr 70  $\phantom{000}70\phantom{000}$ Phe S 8

Gly Pro Val Arg Val Ser Leu Tyr Asp Leu Ala Ser Val Asp Ser Cys

Ser Pro Glu Glu Asn Ser Val Leu Glu Ile Ile Ala Phe His Cys Lys 100 S

His Arg His Arg Met Val Val Leu Glu Pro Leu Asn Lys Leu Leu Gln 115

Leu Asn Phe Leu Cys 140 Phe Lys Trp Asp Leu Leu Ile Pro Lys Phe 130 Ala 2

Asn Leu lle Tyr Met Phe Ile Phe Thr Ala Val Ala Tyr His Gln Pro 160 165 Leu Lys Lys Gln Ala Ala Pro His Leu Lys Ala Glu Val Gly Asn 175 볼 15

Leu Leu Thr Gly His Ile Leu Ile Leu Leu Gly Gly Ile Tyr  $$180\$ Met Ser

2

Leu Leu Val Gly Gln Leu Trp Tyr Phe Trp Arg Arg His Val Phe Ile 195 Trp 11e Ser Phe Ile Asp Ser Tyr Phe Glu Ile Leu Phe Leu Phe Gln 210 210 23

Cys Phe Leu Xaa Ile Glu 235 Ala Leu Leu Thr Val Val Ser Gln Val Leu 235 2

Leu Asn 255 Leu Leu Tyr Thr Arg Gly Phe Gln His Thr Gly 11e Tyr Ser Val 260 Trp Tyr Leu Pro Leu Leu Val Ser Ala Leu Val Leu Gly Trp \$250\$33

Met Ile Gln Lys Pro Trp Xaa 275

**\$** 

(2) INFORMATION FOR SEQ ID NO: 494:

(xi) SEQUENCE DESCRIPTION: SEQ.ID NO: 494: (A) LENGTH: 193 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear 45

Met Ile Arg Cys Gly Leu Ala Cys Glu Arg Cys Arg Trp Ile Leu Pro 1 5 10 15 Arg Gly Trp Leu Gln Ser Ser Asp His Gly Gln Thr Ser Ser Leu Trp 35 Leu Leu Leu Ser Ala Ile Ala Phe Asp Ile Ile Ala Leu Ala Gly 25 20 55

Trp Lys Cys Ser Gln Glu Gly Gly Gly Ser Gly Ser Tyr Glu Glu Gly 8

	WO 98/39448
655	

8 25 20 2 5 S Cys Gln Ser Leu Met Glu Tyr Ala Trp Gly Arg Ala Ala Ala Ala Met 65 70 75 80 ₹ Phe A]a (2) INFORMATION FOR SEQ ID NO: 495: Ala Leu S 8

Leu Phe Cys Gly Phe Ile Ile Leu Val Ile Cys Phe Ile Leu Ser Phe 90 95 Val Thr Tyr Ile Tyr Asn Trp Ala Tyr Gly Fhe Gly Trp Ala Ala Thr 145 Gly Leu Leu Ala Leu Ala Ala Val Phe Gln Ile Ile Ser Leu Val Ile 115 120 125 Glu Asp Asp Leu Leu Gly Asn Ala Lys Pro Arg Tyr Phe Tyr Thr Ser 180 Ile Ile Leu Ile Gly Cys Ala Phe Phe Phe Cys Cys Leu Pro Asn Tyr 165 170 175 Pro Val Lys Tyr Thr Gln Thr Phe Thr Leu His Ala Asn Xaa Ala 130 140 Ξ SEQUENCE CHARACTERISTICS:

Cys Gly Pro Gln Met Leu Val Phe Leu Arg Val Ile Gly 100 105 110

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(2) INFORMATION FOR SEQ ID NO: 496:

SEQUENCE CHARACTERISTICS

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Arg Lys Gln Ala Ser Pro His Arg Ile Leu Phe His Xaa 195 200 205

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Gln Ser Leu Phe Cys Lys Ser Glu Leu Trp Trp Arg Gln Met Arg Ser 165 170 175

Ile Thr Trp Val Pro Ser Pro Lys Ala Gly Trp Arg Trp Thr Lys Gly 185 190

Pro Phe Ser Pro Pro Ala Cys His Thr Ala Pro Asn Ser Val Leu Ile 145 150 150 155

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Pro Pro Pro Asp Ser Asp Leu Cys Ser Gly Pro Leu Leu Pro Gly 130 140

ě Met Ala Leu Thr Leu Leu Pro Ser Val Ser Arg Leu Pro Gly Glu Arg Va. Ser Val Ser Gly Lys Ser Trp Val Gln Val His Tyr Leu Pro Ile Ser Ser Ser Ile Val Asn Tyr Gly Thr 65 70 75 80 Agn Met Lys Val Ile Phe Phe Pro Tyr Pro Val Leu Pro Leu Pro Ala Pro  $35 \hspace{1cm} 40 \hspace{1cm} 45$ Met Ala Ala Ser Gly Leu Pro Tyr Val Leu His His Lys Ser Ser Leu
20 25 30 才 l Glu Leu Pro 115 Pro Gly Leu Ser Gln Leu Pro Thr Ser His Lys Pro Ile Lys Gln 130 135 Pro Thr Gly Thr Trp Val Pro Arg Leu Val Leu Gly Leu Gly Ser Gly Asp 50 55 Ĕ Trp Ser Thr Arg Cys Phe Gln Val Trp Asp Leu Leu Ser 100 105 SEQUENCE DESCRIPTION: SEQ ID NO: 496: (A) LENGTH: 147 amino (B) TYPE: amino acid (D) TOPOLOGY: linear Asp Lys Gly Glu Gly Asn Thr Arg Arg Ala Ser Gly 120 125 Phe acids Leu Val Tyr Pro Leu His Pro 90 95

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SEQUENCE DESCRIPTION: SEQ ID NO: 495:

(D) TOPOLOGY: linear (A) LENGTH: 205 amino acids(B) TYPE: amino acid

Met Ala Ala Gly Asp Gln Val Phe Ser Gly Ala Gly His Val Xaa Glu 1 15

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Xaa Pro Gly Pro Ser Pro Trp Arg Ser Pro Ala Arg Arg Pro Ala Gln
115 120 125

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Arg Gly Pro Thr Trp Thr Ser Pro Pro Thr Arg Val Leu Leu Gly Thr 105 110

Ala Ala Pro Thr Pro Ala Arg Ala Pro Ala Ala Ala Arg Thr Pro Ala 90 95

Leu Trp Met Ala Pro Ala Ala Ala Pro Thr 65 70

Pro Ala Arg Ala Pro Glu 75

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Met Thr Arg Leu Met Arg Xaa Arg Thr Ala Ser Gly Ser Ser Val Ile 50 55

Cys Pro Ala Asn Arg Leu Ser Leu Val Pro Leu Val Pro Ser Ala Ser 35  $40\,$ 

His Val Ala Gly Gly Arg His Ala Trp Leu Leu Thr Trp Gln Ser Ala 25 30

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Glu Tyr Xaa 145

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(2) INFORMATION FOR SEQ ID NO: 497:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 64 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 497:

Met Val Trp Val Leu Trp Ser Ala Pro Ser Leu Ala Pro Pro Trp Val 1 5

9

Gly Pro Cys Trp Pro Ser Thr Gly Asn Cys Cys Leu Cys Glu Val Gly 20 25

Ala Ala Leu Pro Pro Arg Gly Pro Ser Leu Ser Asp Cys Leu Gly. Leu

2

Pro Pro Trp Thr Pro Trp Gly Pro Ala Trp Thr Leu Ala Gln Ser Xaa 50 60

2

22

(2) INPORMATION FOR SEQ ID NO: 498:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 94 amino acids

(B) TYPE: amino acid

30

(D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 498:

Met Ser Thr Gly Ala Leu Asn Thr Ser Pro Pro Ala Ser Asn Arg Leu

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Glu Ser Thr Leu Asn Glu Tyr Leu Ile Gln Pro Gln Leu His Cys Ser

Ser Val Gln Arg Leu Thr Leu Lys Trp Gly Cys Ser Ser Leu Gln Arg \$ 6

Asp Gly Gln Ala Val Pro Trp Gly Leu Trp Gln Arg Ala Tyr Pro Ser 50

5

, 등 Leu Pro Ser Asp Leu Leu Arg Pro His Ala Val 70 Leu Leu Pro Thr

Xaa Ser Val His Thr Cys Glu Ser Ser Pro Ser Val Ser Val တ္တ

(2) INFORMATION FOR SEQ ID NO: 499: 55

(A) LENGTH: 22 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 499: (D) TOPOLOGY: 1inear

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Met Phe Leu Ile Phe Val Tyr Phe Leu Lys Xaa Leu Phe Ser Ser

Leu Pro Phe Leu Trp Leu 20

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(2) INFORMATION FOR SEQ ID NO: 500: 2

(A) LENGTH: 33 amino acids (i) SEQUENCE CHARACTERISTICS:

TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 500: (D) TOPOLOGY: linear

13

Arg Gly Gly Leu Cys Pro Leu Leu Val Pro Gly Pro Leu Ala Arg Gln

Glu Pro Ser Pro Ser Leu Gln Gly Cys Ser Glu Ser Pro Val Gly Met 8

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(2) INFORMATION FOR SEQ ID NO: 501:

(i) SEQUENCE CHARACTERISTICS:

8

(A) LENGTH: 28 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear

35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 501:

Met Gln Phe Leu Leu Thr Ala Phe Leu Leu Val Pro Leu Leu Ala Leu

Cys Asp Val Pro Ile Ser Leu Gly Phe Ser Pro Ser 25

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(2) INFORMATION FOR SEQ ID NO: 502:

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(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 15 andno acids
(B) TYPE: amino acid

(D) TOPOLOGY: linear

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SEQUENCE DESCRIPTION: SEQ ID NO: 502: Œ

Pro Gly Lys Pro Gln Ala Cys Pro Glu Leu Thr Ser Val Leu Pro 1 10 11

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(2) INFORMATION FOR SEQ ID NO: 503:

(1) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 506: 660

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Met Leu Trp Phe Gly Gly Cys Ser Ala Val Asn Ala Thr Gly His Leu 1 5 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 507: (A) LENGTH: 207 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

15

(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 507:

Leu Pro Leu Ala Glu Leu Lys Asn Trp Val 1 5

3 INFORMATION FOR SEQ ID NO: 508:

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(A) LENGTH: 36 amino acids

Ser His Ala Ala Arg Arg His Gln Arg Xaa Leu Leu Ala Ala Ile Asn 115 120 125

Ala Phe Arg Gin Val Arg Leu Lys His Arg Lys Leu Arg Glu Gin Val 130 135

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Gln Gln Asn Leu Ser Ser His Arg Ala Leu Glu Lys Gln Ile Asp 165 170 175

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Thr Leu Ala Gly Lys Leu Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala 180 185

Leu Gly Pro Arg Gln Leu Pro Glu Pro Ser Gln Gln Ser Lys Xaa 195 200 205

Val

35

Leu Gln Glu Ala Trp Met Phe Tyr Lys His Thr Arg Arg Lys Glu 100 105 110

Met Met Asp Ile Gln Tyr Thr Lys Glu Met Lys Glu Ser Ala Ala Arg 85 90 95

Ala Arg Lys Leu Glu Phe Asn Lys Ala Glu Lys His Val His Asn Phe 65 70 75 80

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Ala Gly Lys Leu Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala Leu Gly . 45

Pro Ser Ser Phe Gln Asn Pro Ala Ser Ser Pro Ser Ser Trp Thr His  $50\,$   $^{\circ}$ 

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Asn Leu Ser

Ser Ser His Arg Ala Leu Glu Lys Gln Ile Asp Thr Leu 20 25 30

Met Val Asp Ile Ser Lys Met His Met Ile Leu Tyr Asp Leu Gln Gln 1 . 15

35

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SEQUENCE DESCRIPTION: SEQ ID NO: 505:

30

(2) INFORMATION FOR SEQ ID NO: 505:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 75 amino acids
(B) TYPE: amino acid (D) TOPOLOGY: linear

25

Ser Ile Leu Asn Leu Phe Leu Leu Lys Met Ile Val Ser 20

Met Glu Lys Thr His Arg Leu Arg Ile Arg Asn Pro Cys Leu Gln Phe 1 15

20

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SEQUENCE DESCRIPTION: SEQ ID NO: 504:

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Thr Gly Val Met Gly Val Cys Cys Thr Ala Leu Leu Val Ala Val Val 50 55

Gly Asp Val Val Pro Gly Thr Met Trp Gly Lys Ile Val Cys Leu Cys 35 40 45

Ser Asp Thr Leu Trp Leu Ile Pro Ile Thr Phe Leu Thr Ile Gly Tyr  $20 \ 25 \ 30$ 

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(2) INFORMATION FOR SEQ ID NO: 504:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 29 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Cys Xaa Phe

Asn Lys Ser Leu Xaa Ser Cys Leu Phe Val Leu His Phe Val Leu His 1 15

<u>E</u>.

SEQUENCE DESCRIPTION: SEQ ID NO: 503:

(A) LENGTH: 19 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(1) SEQUENCE CHARACTERISTICS:

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(2) INFORMATION FOR SEQ ID NO: 506:

SEQUENCE CHARACTERISTICS:

(A) LENGTH: 10 amino acids(B) TYPE; amino acid(D) TOPOLOGY: linear

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Glu Glu Glu Pro Gly Tyr Phe Pro Gln Tyr Xaa 65 70 75

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(B) TYPE: amino acid (D) TOPOLOGY: linear

Met Trp Arg Cys Arg Gly Lys Leu Ser Phe Pro Leu Phe Ala Val Val 1 5 10 15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 508:

Ile Val Ser Cys Arg Lys Asp Gly Pro Asp Ala Ala Ala Ala Pro Ala

Val Xaa Lys Lys 35

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(2) INFORMATION FOR SEQ ID NO: 509:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 19 amino acids

(B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 509: (D) TOPOLOGY: linear

Met Ala Leu Val Ala Leu Phe Thr Gln Leu Met Arg Xaa Leu Gly Arg 22

Cys Pro Gln

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(2) INFORMATION FOR SEQ ID NO: 510:

(A) LENGTH: 32 amino acids (1) SEQUENCE CHARACTERISTICS:

35

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 510:

6

Phe Asp Ser Trp Arg Glu Gln Thr Arg Thr Asn Ile Gln Pro Gln Arg

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(2) INFORMATION FOR SEQ ID NO: 511: 20

(A) LENGTH: 28 amino acids SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 511:

Met His Leu Leu Asp Phe Phe Arg Asp Leu Val Leu Leu Val Leu 115  $^{\rm 15}$ 

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Ala Leu Leu Asp Ser Phe Trp Leu Glu Val Gln Lys 20

(2) INFORMATION FOR SEQ ID NO: 512:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 26 amino acids (B) TYPE: amino acid

2

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 512:

Met Cys Leu Ile His Phe Ile Lys Ile Ile Leu Val Phe Ile Leu Lys

13

Leu Trp Leu Tyr Ser Gln Lys Cys Pro Lys 20 25

2

(2) INFORMATION FOR SEQ ID NO: 513:

(i) SEQUENCE CHARACTERISTICS:

25

(A) LENGTH: 33 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 513:

30

Met Ile His Val His Glu Trp Asn Asp Gln Met Leu Met Val Tyr Ile

Phe Leu Tyr Pro Val Ser Ile Thr Phe Leu Asn Leu Cys Ser Leu Thr 8 35

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(2) INFORMATION FOR SEQ ID NO: 514:

(i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 514: (A) LEWSTH: 47 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

Leu Asn Glu Ser Tyr Val Ser Arg Ala Gly Gly Trp Phe Ser Met Phe

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Xaa Leu Ile Phe Phe Leu Leu Ala Leu Gly Ser Xaa Leu Cys Leu Leu 20

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Leu Cys Leu Pro Ser Phe Asn Lys Thr Arg Arg Lys Gln Lys Pro

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3 40 35 છ 23 8 2 5 55 Š 8 Leu Asn Trp 1 Leu (2) INFORMATION FOR SEQ ID NO: 515: Ser Ser Lys Thr Pro Leu Pro Ser Glu Arg Arg Trp Ile Ser Gly Ser 1 10 15 Leu Asp Leu His Lys Ser Leu Val Cys Thr Ala Leu Arg Gly Lys Leu 50 55 60 Val Asp Ser Gln Met Asp Asp Met Asp Met Asp Leu Asp Lys Glu Phe 20 30 Phe Ala Phe Cys Ala Glu Leu Met Ile Gln Asn Trp Thr Leu Gly Ala 1 10 15 (2) INFORMATION FOR SEQ ID NO: 516: Leu Phe Val Asp Leu Val Glu Lys Phe Val Glu Pro Cys Arg Ser Asp 100 105 Gly Val Phe Ser Glu Met Glu Ala Asn Phe Lys Asn Leu Ser Arg Gly 65 70 75 80 Leu Gln Asp Leu Lys Glu Leu Lys Val Leu Val Ala Asp Lys Asp Leu 35  $40\,$ (2) INFORMATION FOR SEQ ID NO: 517: Leu Val ij rea (1) SEQUENCE CHARACTERISTICS: ž Ala Leu Glu Pro Ala Leu Gly His Trp 35 40 Met Ala Pro Arg Pro Trp Leu Leu Gly Ile Ala Leu Leu Gly 20  $25\,$  30 Asn Val Ala Ala Lys Leu Thr His Asn Lys Asp Val Arg Asp  $90 \ 95$ E χ (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 515: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 517: SEQUENCE DESCRIPTION: SEQ ID NO: 516: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 43 amino acids (A) LENGTH: 174 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 3 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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His Trp Pro Leu Ser Asp Val Arg Phe Phe Leu Asn Gln Tyr Ser Ala 115 120 125

Thr Trp Ala Pro Ser Ala Ala Ala Ser Cys Ala Cys Ile Met Thr Glu 145 150 Ser Val His Ser Leu Asp Gly Phe Arg His Gln Ala Ser Gly Thr Ala 130 135 140

S

5 Val Pro Pro Asn Ala Pro Pro Thr Leu Thr Ile Lys Leu Leu
165 170

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(2) INFORMATION FOR SEQ ID NO: 518:

(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 43 amino acids

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 518: (B) TYPE: amino acid
(D) TOPOLOGY: linear

25 Met Trp Lys Asn Leu Gly Ser Gly Ser Val Phe Val Thr Trp Phe Ser 1 15 Leu Val Met Ile Leu Ser Gly Ile Gly Pro Leu Gly Asp Ala Glu Asp 20 25 30

30 Ser Ile Ser Asp Val Ser His Arg Leu Arg Pro 35 40

35 (2) INFORMATION FOR SEQ ID NO: 519:

(1) SEQUENCE CHARACTERISTICS: Ĕ SEQUENCE DESCRIPTION: SEQ ID NO: 519: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 13 amino acids

6

25 Phe Gln Phe Pro Leu Leu Thr Ile Ala Leu Gln Phe Leu 1 5 10

(2) INFORMATION FOR SEQ ID NO: 520

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 30 amino acids(B) TYPB: amino acid(D) TOPOLOGY: linear

50

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 520:

55

Met His Tyr Val Ile Val Leu Ser Leu Phe Val Val Leu Glu Lys Lys
1 10 15

Asn Lys Met Gly Ser Asp Gly Cys Leu Arg Lys Asn Gly Ser 20 25 30

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(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 47 amino acids (B) TYPE: amino acid (2) INFORMATION FOR SEQ ID NO: 521: S

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 521: (D) TOPOLOGY: linear

2

Met Ser Arg Ser Ile Val Leu Arg Gly Ser Leu Phe Leu Phe Ser

His Tyr Thr Leu Lys Leu Leu Ser Val Ile Lys Gln Thr Asn Arg Lys 13

Ile Val Trp Glu Lys Pro Cys Ile Arg Leu Phe Tyr Xaa Val Leu 35 46

2

(2) INFORMATION FOR SEQ ID NO: 522

(A) LENGTH: 26 amino acids (i) SEQUENCE CHARACTERISTICS: 23

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 522;

3

Pro Leu Pro Val Leu Leu Cys Leu Thr Leu Pro Met Pro Leu Pro 10 Met

Ser Ala Thr Ala Arg Gly Gly Asn Arg Thr 20 25

35

(2) INPORMATION FOR SEQ ID NO: 523:

9

(A) LENGTH: 58 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid 3

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 523:

(D) TOPOLOGY: linear

45

Ser Ser Ile Pro Val Ser Ile Leu Ile Gly Met Lys Leu Ile Leu Tyr

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Leu Leu Ile Thr Glu Ser Gly Ser His Glu Lys Lys Ser Phe Tyr Pro 20 30 Ser Phe Lys Tyr Met Phe Lys Ile Ile Ile Tyr Val Ser Ala Tyr Cys 35

Arg Thr Ala Leu Arg Ala Thr Val Ser His

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(2) INFORMATION FOR SEQ ID NO: 524:

999

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 19 amino acids

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 524:

Asn Arg Thr Leu Leu Phe Leu Ile Leu Phe Val Leu Phe Gly Leu Gly

Tyr Gly Phe

9

15

(2) INFORMATION FOR SEQ ID NO: 525:

(A) LENGTH: 40 amino acids (1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid

2

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 525: (D) TOPOLOGY: linear

Phe Leu Leu Val Leu Ser Val Phe Cys Asp Phe Met Cys Ser Ile  $_{\rm 5}$ Met 25

Ala Pro Arg Cys His Ala Leu Ser Leu Val Ser Leu Arg Ala Gln His  $25\,$ 

Leu Phe Ile Thr Cys His 35 Ser Leu

2

(i) SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 526:

35

(A) LENGTH: 57 amino acids (B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 526:

Met Leu Leu Phe Ile Leu Leu Thr Leu Ser Ser Gly Cys Arg Leu Leu  $1 \ \ \, 1$ 

45

Val Ser Ser Trp Lys Thr Phe Leu Pro His Phe Ser Leu Pro Gly Pro 20 30

Arg Glu His Pro Glu Gly Ser Arg Thr Trp Phe Phe Arg Tyr Trp Glu 35 S

Pro Gly Ala His Cys Leu His Cys Ala 50 55

23

(2) INFORMATION FOR SEQ ID NO: 527:

(1) SEQUENCE CHARACTERISTICS:

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(A) LENGTH: 21 amino acids (B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 527: (D) TOPOLOGY: linear

S

Ala Arg Leu Leu Phe Leu Ser Ser Val His Pro Ser Ile Met Pro 1 15

Ser Cys Asn Gln Leu 20

5

(2) INFORMATION FOR SEQ ID NO: 528:

(i) SEQUENCE CHARACTERISTICS:

15

(A) LENGTH: 39 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 528:

20

Met Ser Leu Thr Ser Ser Leu Thr Phe Leu Ser His Ile Leu Leu Leu 1 5 10 (X

Pro Gln Lys Leu Gln Phe Leu Ser  $\mathfrak{T}\text{rp}$  Met Glu Arg Gln Gln Arg Cys 25

25

Trp Gly Phe Gly Xaa Lys Thr Ala Ser Phe Gly Ala Val Gly Glu Thr 35

Leu Gly Gly Arg Cys Ile Gln Gly Arg Phe Ala Ser His Ser Lys Phe
20 25 30

Pro

Pro Asp Gln Glu Pro Gln Lys Glu Thr Glu Pro Ala Thr Ser Ser 50 55

20

Met Gly Ser Ser Val Leu Pro Phe Cys Val Cys Val Thr Ser Pro Ser

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SEQUENCE DESCRIPTION: SEQ ID NO: 530:

(A) LENGTH: 82 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 530:

(i) SEQUENCE CHARACTERISTICS:

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Thr Gly Val Ala Lys Tyr Ala

8 His Ala Arg Pro Trp Ala Arg Val Ile Gly Leu Arg Ile Trp Pro Gln 65 70 75

Pro Asn

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2 INFORMATION FOR SEQ ID NO: 531:

Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 20 amino acids

8

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 531: (B) TYPE: amino acid
(D) TOPOLOGY: linear

45

Met Leu Leu Ser Val Ala Ile Phe Ile Leu Leu Thr Leu Val Tyr Ala
1 10 15

Ş Tyr Trp Thr Met 20

(2) INFORMATION FOR SEQ ID NO: 532:

S

E SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 75 amino acids

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 532: (D) TOPOLOGY: linear

8

8

Ser Pro Gly Leu Lys Asn'Gly Ile Phe Leu Phe Leu Leu Arg Ala Met 100 105 110 .

S

Trp Val

Arg Val Ile His Pro Phe Leu Val Leu Pro 85

o Cys Leu Tyr 95

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His Trp Val His Gly His Leu Pro Trp Cys His Pro Tyr Ile Gln Val 50 55

Glu Phe Ser Ala Leu Ile Glu Ser Thr Ala Gln Leu Gly Leu Pro Phe 65 70 75 80

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SEQUENCE DESCRIPTION: SEQ ID NO: 529:

(A) LENGTH: 128 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Val Leu Arg Leu Ile Gln Leu Ile Phe Leu Ile Phe Phe Ile His 1 5 10

Ile Ile Leu Leu Ile Pro Gly Ser Arg Pro Cys Gly Ser Trp Val 20

Asn Asp Arg Xaa Leu Gly Leu Arg Asp Val Thr His Leu Ile Tyr Leu 35  $40\,$ 

35

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SEQUENCE CHARACTERISTICS:

(2) INFORMATION FOR SEQ ID NO: 529:

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Pro Gly Gly Met Phe Pro Gly Asn Leu Glu Ala Phe Arg Val Pro Val 115 120 120

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Asn Cys Glu Ile Leu Glu Tyr Cys Tyr Tyr Leu Thr Gln Leu Lys Ile Arg Cys Ser Ile Thr Ala Val Ser Asp Ser Ser Thr Ser Trp Ala Ile Ser Met Gly Lys Tyr Leu Ser Ile Pro Thr Val Leu Leu Lys Ile Ile Lys Ala Gln Leu Lys Ile Glu Asn Lys Asp Leu Asp Asn Lys Thr Ala 50 Lys Gly Gly Gly Gln Glu Ala Leu Thr Cys Thr 65 75 15 9

(2) INFORMATION FOR SEQ ID NO: 533: ន

SEQUENCE DESCRIPTION: SEQ ID NO: 533: (A) LENGTH: 60 amino acids (1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear Ī 23

Met Phe Leu Met Arg Met His Leu Cys Phe Cys Lys Tyr Cys Cys Ser  $1 \\ 1$ Phe Ile Val Thr Pro Thr Ser Thr Ser Asn Thr Xaa Ser Tyr Leu Trp 20 9

Pro Trp Ile Ser Ala Ser Met Ala Gly Arg Gly Ser Xaa Trp Ala Cys Thr Leu Asn Ala Val Thr Arg Glu Gly Leu Pro Glu
50 55 60 35

(2) INFORMATION FOR SEQ ID NO: 534: 6

(A) LENGTH: 39 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: 1inear

45

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 534:

Ser Leu Leu Asn Thr His Thr Leu Cys Phe Val Leu Phe Cys Phe 5 10 Thr Leu Ser Ile Asn Gln Glu Lys Leu Ala Asn His Leu Ala Phe Arg 20 ႙

Ile Leu Phe Phe Ile Val Phe 35 55

(2) INFORMATION FOR SEQ ID NO: 535: 8

(A) LENGTH: 2 amino acids SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 535:

Met Leu

9

(2) INFORMATION FOR SEQ ID NO: 536:

(A) LENGTH: 36 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

(i) SEQUENCE CHARACTERISTICS:

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 536:

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Met Asp Gln Phe Lys Ile Phe Tyr Phe Leu Lys Ala Phe Phe Ala Cys 1 5

Cys Asn Val Gln Asp Pro Ser Pro Phe Met Gly Glu Thr Gly Ser Tyr 20 20

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Leu Asn Ile Gly

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(2) INFORMATION FOR SEQ ID NO: 537:

(A) LENGTH: 14 amino acids (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 537:

Met Phe Asp Phe Leu Ser Tyr Phe Lys Asp Leu Leu Ser Cys  $_1$ 6

(2) INFORMATION FOR SEQ ID NO: 538: 45 (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 18 amino acids TYPE: amino acid (D) TOPOLOGY: linear

S

Met Gly Phe Gly Phe Val Leu Asn Ile Phe Ser Phe Phe Leu Xaa Pro

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 538:

Pro Leu

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5 55 30 20 5 8 50 8 ઝ 25 3 Leu Leu Tep Thr Leu Leu Ala Xaa Tyr Xaa 1 5 10 (2) INFORMATION FOR SEQ ID NO: 539: Phe Val Leu Gln Arg Leu Val Arg Asn Val Glu Tyr Tyr Gln Ser Asn Tyr Val Phe 65 70 75 Glu Trp Leu Glu Arg Arg Arg Ala Thr Ile Arg Pro Trp Ser Thr Phe \$45\$Ser Gly Thr Thr Leu Leu Pro Lys Leu Ile Pro Ser Gly Ala Gly Arg  $20 \ \ 30$ Met Ala Ala Gin Lys Asp Gin Gin Lys Asp Ala Giu Ala Giu Gly Leu 1 15 (2) INFORMATION FOR SEQ ID NO: 540: (2) INFORMATION FOR SEQ ID NO: 541: Val Phe Leu Gly Leu Ile Leu Tyr Cys Val Val Thr Ser Pro Met Leu 95 Leu Arg Thr Leu Glu Ser Lys Leu Val Leu Phe Gly Arg Glu Val Ser 35 40 Val Ala Leu Ala Val Phe Phe Gly Ala Cys Xaa 100 105 Leu Val (i) SEQUENCE CHARACTERISTICS: Ě Ě Phe Leu Gly Leu Ile Leu Tyr Cys Val Val Thr Ser Pro Met
15 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 541: . E SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS SEQUENCE DESCRIPTION: SEQ ID NO: 539: SEQUENCE DESCRIPTION: SEQ ID NO: 540: Ala Leu Ala Val Phe Phe Gly Ala Cys Tyr Ile Leu Tyr 20 25 30 (A) LENGTH: 108 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 11 amino acids (A) LENGTH: 106 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

> 5 S Pro Ala His Gln Tyr Ala Leu Ala Gly Gly Ile Ser Phe Pro Phe Phe 50  $\,$  50  $\,$ Asp Gly Glu Glu Leu Gln Met Glu Pro Val Leu Val Val Ile Gly Ser His Ala Ala Phe His Gln Ile Glu Ala Val 95 95 Trp Leu Ala Gly Ala Gly Ser Ala Val Phe Trp Val Leu Gly Ala Thr  $65 \qquad 70 \qquad 75 \qquad 80$

2 (2) INFORMATION FOR SEQ ID NO: 542:

30 23 20 Ser Phe Asn Ile Gly Asp Ser Ser Ser Gly Leu Ile Gln Thr Val Phe Ile 20 25 Met Asp Arg Phe Thr Val Ala Gly Val Leu Pro Asp Ile Glu Gln Phe 1 15 Ser (i) SEQUENCE CHARACTERISTICS Tyr Met Val Leu Ala Pro Val Phe Gly Tyr Leu Gly Asp Arg 35 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 542: (A) LENGTH: 136 amino acids
(B) TYPE: amino acid

Ala Thr Leu Gln Ala Pro Lys Xaa 130 135 Gly Cys Leu Leu Thr Arg Gly Leu Val Gly Val Gly Glu Ala Ser Tyr Ser Thr 95Val Thr Leu Gly Ser 65 Tyr Asn Arg Lys Tyr Leu Met Cys Gly Gly Ile Ala Phe Trp Ser Leu 50 60 Ile Ala Pro Thr Leu Ile Ala Asp Leu Phe Val Ala Asp Gln Arg Thr 100 105 Ser Ala Ser Ser Thr Leu Pro Phe Arg Trp Ala Val Val Trp 115 120 128 Ser Phe Ile Pro Gly Glu His Phe Trp Leu Leu 70 75 80

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INFORMATION FOR SEQ ID NO: 543:

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(1) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 543: (A) LENGTH: 424 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Pro Pro Arg 30 Leu Ala Ser Arg Val Val Leu Gly Glu 90 95 Met Ala Gly Asp Trp His Trp Ala Leu Arg Val Thr Pro Gly Leu Gly 1 5 Ser Ala Asp Leu Arg Ala Leu Ala Arg Asn Pro Ser Phe Val Leu 55 ile Phe Gly Leu Ile Thr Cys Leu Thr Gly Val Leu Gly Val Gly Leu 115 Gly Val Glu Ile Ser Arg Arg Xaa Arg His Ser Asn Pro Arg Ala Asp 130 Thr Gly Leu Leu Gly Ser Ala Pro Phe Leu Phe 150 Ser Ile Val Ala Thr Tyr Ile Phe 170 Ser Met Asn Trp Ala Ile Val Ala 185 Ser Leu Leu fyr Val Val Ile Pro Thr Arg Arg Ser Thr Ala Glu 205 Ala Phe Gin Ile Val Leu Sar His Leu Leu Gly Asp Ala Gly Ser Pro 210 Tyr Leu Ile Gly Leu Ile Ser Asp Arg Leu Arg Arg Asn Trp Pro Pro 225 Ala Ala Pro Pro Ala Cys Pro Trp Pro Val Cys Ser Ser Glu Arg Leu 320 Leu Ser Glu Phe Arg Ala Leu Gln Phe Ser Leu Met Leu Cys 245 255 Gly Ala Leu Gly Gly Ala Leu Ser Trp Ala Pro Xaa Ser 260 260 Ser Leu Arg Pro Thr Ala Gly Gly His Ser Cys Thr Cys Arg Ala Cys 275 Lys Gln Gly Pro Gln Thr Thr Gly Leu Trp Cys Pro Ser Gly 295 300 Ser Asp Leu Pro Pro Leu Asn Pro Thr 40 45 Thr Ale Val Ala Phe Val Thr Gly Ser 70 The Pro Pro Cys Leu Pro Gly Asp Ser Cys Ser Ser Asp 100 110 Leu Val Val Arg Glu 25 Phe Leu Leu Arg P P Leu Ala Cys Ala Arg Gly ž Z Phe Ile Gly Glu Thr Leu 180 Ala Val Glu Arg His 35 Ę Leu. Phe Leu Trp Ala Pro Ala 85 Leu Val Cys Ala Val Val Ala Val Leu Gly Leu 195 Ala Phe Val £ 8 Ser 7 290 Leu Ser Asp 11e Ser Phe Pro 145 Ile ŢŢ 65 ž S 으 15 ຊ 22 33 3 <del>6</del> 5 8 55

Pro Leu Thr Tyr Leu His Ile Cys His Ser Xaa Pro Trp Ala His Pro 335

Thr Lys Gly Leu Gly Leu Thr Pro Trp Pro Gly Pro Ala Ser Arg Gly 346

Ser Gly Ser Thr Leu Gly Arg Val Pro Ala Pro Arg His Tyr Xaa Gly 365 365 Glu Glu Val Gly Val Gln Glu Gly Asp Pro Ser Pro Gln Gly Xaa Pro 370 380 Gin Gly Leu Gly Ala Ile Cys Asn Gly Ile Lys Phe Val Ala Arg Pro 395 395 2

Gin Val Pro Ala Leu Val Phe Leu Trp Val Ala Ser Asp Leu Ala Pro

2

Leu His Pro Arg Ala Pro Glu Arg

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(2) INFORMATION FOR SEQ ID NO: 544

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(A) LENGTH: 39 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

Phe Arg Phe Val Ile Cys Leu Phe Leu Trp Leu Val Leu Cys Arg 10 SEQUENCE DESCRIPTION: SEQ ID NO: 544: (X Met

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Ser Ala Ser Arg Ile Ala Leu Tyr Tyr Arg Ile Val Phe 20 Ser Thr Asp 35

Leu Ile His Gln Cys Ser 35

Ser

<del>4</del>

INFORMATION FOR SEQ'ID NO: 545: 3

45

(A) LENGTH: 58 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (i) SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 545:

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Leu Pro Try Xea Ala Gin Leu Leu Asp Arg Thr Ile Gly Pro Leu 5 Tyr Leu Leu Phe Val Gln Phe Ser Pro Ala Phe Ser Arg Thr Ser Pro 20 Met

Trp Arg Ser Pro Lys Asn Phe Arg Arg Leu Tyr Pro Pro Cys Thr Thr 35

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Ser Gly Cys Ala Ala Arg Trp Leu Phe Ser

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50 (1) SEQUENCE CHARACTERISTICS:

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(2) INFORMATION FOR SEQ ID NO: 546:

(B) TYPE: amino acid (A) LENGTH: 33 amino acids

<u>£</u> SEQUENCE DESCRIPTION: SEQ ID NO: 546:

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Met Gly Leu Ser Val Leu Leu Pro Leu Cys Leu Leu Gly Pro Gly Arg 1 10 15

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Phe Thr Ser Gly Gln Lys Pro Leu Asp Thr Pro Gly Leu Gly Val Pro 20 30

Phe

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Ξ SEQUENCE CHARACTERISTICS: (A) LENGTH: 367 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 547:

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 547:

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Met Ala Lys Pro Gin Val Val Val Ala Pro Val Leu Met Ser Lys Leu 1 15

Ser Val Asn Ala Pro Glu Phe Tyr Pro Ser Gly Tyr Ser Ser Ser Tyr 20 30

35

Thr Glu Ser Tyr Glu Asp Gly Cys Glu Asp Tyr Pro Thr Leu Ser Glu 35

Tyr Val Gln Asp Phe Leu Asn His Leu Thr Glu Gln Pro Gly Ser Phe 50 60

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Glu Thr Glu Ile Glu Gln Phe Ala Glu Thr Leu Asn Gly Cys Val Thr .65  $70\,$   $75\,$ Thr Asp Asp Ala Leu Gln Glu Leu Val Glu Leu Ile Tyr Gln Gln Ala 95

25

50 Thr Ser Ile Pro Asn Phe Ser Tyr Met Gly Ala Arg Leu Cys Asn Tyr 100 105

Leu Ser His His Leu Thr Ile Ser Pro Gln Ser Gly Asn Phe Arg Gln
115 120 125

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Ala Lys Gly Asp Glu Vali Thr Arg Lys Arg Phe His Ala Phe Val Leu 145

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Phe Leu Gly Glu Leu Tyr Leu Asn Leu Glu Ile Lys Gly Thr Asn Gly
175

Gln Val Thr Arg Ala Asp Ile Leu Gln Val Gly Leu Arg Glu Leu Leu 180

Asn Ala Leu Phe Ser Asn Pro Met Asp Asp Asn Leu Ile Cys Ala Val 200 205

Lys Leu Leu Lys Leu Thr Gly Ser Val Leu Glu Asp Ala Trp Lys Glu 210 225

5

5 Lys Gly Lys Met Asp Met Glu Glu Ile Ile Gln Arg Ile Glu Asn Vel 225 230 230

Val Leu Asp Ala Asn Cys Ser Arg Asp Val Lys Gln Met Leu Leu Lys 245 250 255

8 Leu Val Glu Leu Arg Ser Ser Asn Trp Gly Arg Val His Ala Thr Ser 260 265 270

25 Glu Pro Thr Phe Tyr Thr Ser Asp Gly Val Pro Phe Thr Ala Ala Asp 290 300 Thr Tyr Arg Glu Ala Thr Pro Glu Asn Asp Pro Asn Tyr Phe Met Asn 275 280 285

ઝ Pro Asp Tyr Gln Glu Lys Tyr Gln Glu Leu Leu Glu Arg Glu Asp Phe 105 310 315 Phe Pro Asp Tyr Glu Glu Asn Gly Thr 325 Asp Leu Ser Gly Ala Gly Asp 330

35 Pro Tyr Leu Asp Asp Ile Asp Asp Glu Met Asp Pro Glu Ile Glu Glu 340 345 350

Ala Tyz Glu Lys Phe Cys Leu Glu Ser Glu Arg Lys Arg Lys Gln 355

6

(2) INFORMATION FOR SEQ ID NO: 548:

E SEQUENCE CHARACTERISTICS

45

(A) LENGTH: 77 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 548:

50

Met Leu Arg Leu Asp Ile Ile Asn Ser Leu Val Thr Thr Val 1 5 10 Phe Met

Leu Ile Val Ser Val Leu Ala Leu Ile Pro Glu Thr Thr Thr Leu Thr 20 25 30

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Asp Gly Ala Leu Ile Tyr Arg Lys Leu Leu Phe Asn Pro Ser Gly Pro

Val Gly Gly Val Phe Ala Leu Val Thr Ala Val Cys Cys Leu Ala 35 40 45

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Pro Val His Glu Lys Lys Glu Val Leu 70 fyr Gln Lys Lys 65

INFORMATION FOR SEQ ID NO: 549; 2

(A) LEWSTH: 47 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear Ξ 2

. Met Leu Lys Gln Val Met Phe Val Phe Ser Gly Met Gly Pro Arg Ser l  $_{\rm 1}$ 

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 549:

2

His Cys Trp Gly Leu Pro Leu Ala Cys Gly Thr Phe Val Gln Gly His 20 Ser Ser His Leu Leu Pro Leu Lys His Gln Gly Ala **\$** Gln Ala Asp S

2

(2) INFORMATION FOR SEQ ID NO: 550:

25

(A) LENGTH: 168 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 8

Met Leu Leu Ser Leu Ala Ala Phe Ser Val Ile Ser Val Val Ser Tyr Leu lle Leu Ala Leu Leu Ser Val Thr lle Ser Phe Arg lle Tyr Lys \$20\$35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 550

Gln Lys Ser Glu Glu Gly His Pro Phe Lys Ser Val Ile Gln Ala Val 6

Glu Ala Phe His Asn 60 116 3 ŝ ž Asn Arg Ala I 75 Ala Tyr Leu Asp Val Asp Ile Thr Leu Ser Ser 50 Val His Ile Met 70 Met Asn Ala Ala 45

Ile Arg Leu Phe Leu Val Glu Asp Leu Val Asp Ser Leu Lys Leu Ala 95 S

Het Try Leu Met Thr Tyr Val Gly Ala Val Phe Asn Gly Ile 100 Val Phe 55

Thr Leu Leu lle Leu Ala Glu Leu Leu Ile Phe Ser Val Pro Ile Val 120

Tyr Glu Lys Tyr Lys Thr Gln Ile Asp His Tyr Val Gly Ile Ala Arg 130 8

Asp Gln Thr Lys Ser Ile Val Glu Lys Ile Gln Ala Lys Leu Pro Gly 145

s Lys Ala Glu Xaa 165 Ile Ala Lys Lys

S

INFORMATION FOR SEQ ID NO: 551: 3 2

(A) LENGTH: 124 amino acids (B) TYPE: amino acid SEQUENCE CHARACTERISTICS:

SEQUENCE DESCRIPTION: SEQ ID NO: 551: ê Ī

2

Val Pro Phe His Leu Leu Val Val Leu Arg Ser Arg Ala Val Arg 5

Ala Arg Arg Arg Glu Pro Arg Ser Leu Pro Arg Pro Gly Asp Glu 25

2

Glu Leu Gin Leu Leu Cys Gly Ala Arg Ser Asp Phe Leu Glu Arg

23

Cys Glu Glu Asp Trp Val Cys Leu Trp His His Ala Asp His Ala Ala 50 60

Phe Pro Gly Ser Phe Gln Cys His Gln Cys Gly Phe Leu Pro His Pro 65 Gly Ser Ser Leu Cys His His Gln Leu Gln Asp Leu Gln Val Arg His 85 9

Ser Cys Thr Glu Val Arg Arg Pro Ser Ile Gln Ser Leu Pro 100 Pro

35

Š Leu Arg Ser Phe 120 Gly Arg Arg His Tyr Ser Val 115

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(2) INFORMATION FOR SEQ ID NO: 552:

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SEQUENCE DESCRIPTION: SEQ ID NO: 552: Œ

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Met Val His Leu Leu Val Leu Ser Gly Ala Trp Gly Met Gln Met Trp 1 5 10 15 Pro Arg His Val Thr Phe Val Ser Gly Phe Leu Leu Phe Arg Ser Leu 20 55

Thr Phe Gly Leu Val Gln Ser Lys Leu Phe. Pro Phe Tyr Phe His Ile \$ 6

Ser Met Gly Cys Ala Phe Ile Asn Leu Cys Ile Leu Ala Ser Gln His 50 55

S Ala Trp Ala Gln Leu Thr Phe Trp Glu Ala Ser Gln Leu Tyr Leu Leu 65 70 75 Phe Leu Ser Leu Thr Leu Ala Thr Val 85 Asn Ala Arg Trp Leu Glu Pro 90 95

5 Gly Leu Gly Gly Glu Val Pro Gly Ser His Gln Gly Pro Asp Pro Tyr
115
120
125 Arg Thr Thr Ala Ala Met Trp Ala Leu Gln Thr Val Glu Lys Glu Arg  $100\,$ 

5 Arg Gln Leu Arg Glu Lys Asp Pro Lys Tyr Ser Ala Leu Arg Gln Asn  $130\,$ 

20 Phe Phe Arg Tyr His Gly Leu Ser Ser Leu Cys Asn Leu Gly Cys Val 145

Leu Ser Asn Gly Leu Cys Leu Ala Gly Leu Ala Leu Glu Ile Arg Ser 165 170 175

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30 (2) INFORMATION FOR SEQ ID NO: 553:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 72 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

<u>X</u> SEQUENCE DESCRIPTION: SEQ ID NO: 553:

35

Glu Gln Asn Ser Ala Thr Val Glu Pro Ser Ser His Glu Ile Leu His 20 25 30 Met Ala Phe Ile Leu Leu Phe Tyr Cys Leu Met Thr Phe Leu Ser Leu 1 5

6

\$ Leu Gln Asn Cys Phe Glu Leu Leu Arg Thr Ser Thr Ser Gln Cys 35 40

Thr Glu Gly Ile Pro Cys Ala Lys Ile Pro Glu Trp Val Thr His Leu 50 55 60

50 Thr Trp Gln Thr Leu Lys Asn Ser 65 70

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(2) INFORMATION FOR SEQ ID NO: 554:

Ξ (a) LENGTH: 45 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Val Leu Arg Ile Ile Cys Leu Trp Pro Cys Gly Thr Thr Leu Pro Leu 1 15 (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 554:

Val Glu Lys Ala His Asp Ser His Ser Ala Asp Pro Val Cys Pro Gly
25 30

5 Leu Thr Ala His Leu Pro Val Leu Leu Tyr Val Gln Leu 35 40 45

(2) INFORMATION FOR SEQ ID NO: 555:

5

(i) SEQUENCE CHARACTERISTICS (A) LENGTH: 251 amino acids (B) TYPE: amino acid

20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 555: (D) TOPOLOGY: linear

ઝ 25 Gly Phe Met Ile Val Tyr Asn Phe Ser Leu Val Ala Leu Ser Leu Tyr 50 55 Ser Leu Gly Pro Arg Ile Met Ala Asn Arg Lys Pro Phe Gln Leu Arg 35 40 45 Pro Leu Leu Met Thr Ser Ile Leu Leu Thr Tyr Val Tyr Phe Val Leu 20 25 Met Lys His Ala Asp Pro Arg Ile Gln Gly Tyr Pro Leu Met Gly Ser 1 15

35 Arg Cys Asp Pro Gln Asp Cys Thr Leu Gly Gln Cys Pro Ser Val Pro 85 90 95 Ile Val Tyr Glu Phe Leu Met Ser Gly Trp 65 70 Leu Ser Thr Tyr Thr Trp 75

Ser Pro Xaa Thr Pro Val Thr Lys Ala Tyr Val Val Arg Thr Glu Gln 105 110

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Trp Phe Leu Thr His Phe Pro Arg Ala Ala Pro Gly Met Trp Pro His 130 135 Gly Thr Gly Pro Pro Leu Pro Thr Ala Ala Leu Gln Gly Pro Arg Leu 115 120 125

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Cys Cys Leu Pro Leu Gln Ser Trp Gly Leu Lys Gly Leu Tyr Ser Tyr
145 150 150

Phe Pro Leu Pro Ala Leu Lys Leu Gly Arg Gly Ala Leu Arg Ala Gly
175

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Thr Lys Gly Leu Val Ala Phe Phe Leu Thr Gln Lys Arg Ser Ala 180 185

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Ile Met Ser Leu Trp Thr Gln Ser His Ser Ser Thr Pro His Thr Glu
195 200 205

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Ala Val Ala Ser Gly Pro Lys Val Arg Val Gly Gly Gly Leu Gly Ile 210

A50 240 Gin Pro Val Glu Ala Ala Tyr Ser Thr Cys Val Leu Ile Lys Ser 235

Arg Gly Asn His Glu Lys Lys Lys Lys Lys Lys 250

(2) INFORMATION FOR SEQ ID NO: 556:

2

(A) LENGTH: 19 amino acids SEQUENCE CHARACTERISTICS 3

2

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 556:

Leu Ala Gly Leu Cys Gly Gln Leu Ser Ser Pro Ala Leu Cys Val 10 Gly ೫

Asn Arg Leu

25

(2) INFORMATION FOR SEQ ID NO: 557:

(A) LENGTH: 217 amino acids SEQUENCE CHARACTERISTICS:

2

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 557:

35

Met Ile Thr Glu Lys Trp Gly Leu Asn Met Glu Tyr Cys Arg Gly Gln 1 5 15

Val Val Ala 30 Ser Arg Leu Leu Glu Lys Tyr Pro Gln Ala Ile Tyr Thr Leu Cys Ser 35 Ę Ser Lys Met Ala Tyr Ile Xaa Ser Ser Gly Phe Ser 20

6

Trp Leu Ala Lys Ser Val Pro Val Met Gly 55 Ser Cys Ala Leu Asn Met 50 45

Ser Val Ala Leu Gly Thr Ile Glu Glu Val Cys Ser Phe Phe His 70 S &

S

Pro Gln Leu Leu Glu Leu Asp Asn Val Ile Ser Val Leu 95 Arg Ser

Phe Gln Asn Ser Lys Glu Arg Gly Lys Glu Leu Lys Glu Ile Cys His 110 55

Ser Oln Trp Thr Cly Arg His Asp Ala Phe Glu Ile Leu Val Glu Leu 115 Leu Gln Ala Leu Val Leu Cys Leu Asp Gly Ile Asn Ser Asp Thr Asn è

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Ile Arg Trp Asn Asn Tyr Ile Ala Gly Arg Ala Phe Val Leu Cys Ser 145

Phe Asp Phe Ile Val Thr Ile Val Val Leu Lys Asn 165 Ala Val Ser Asp

Val Leu Ser Phe Thr Arg Ala Phe Gly Lys Asn Leu Gln Gly Gln Thr 180

9

Ser Asp Val Phe Phe Ala Ala Gly Ser Leu Thr Ala Val Leu His Ser 195

Leu Asn Glu Val Ile Gly Lys Tyr Xaa 210

15

(2) INFORMATION FOR SEQ ID NO: 558: 8

(i) SEQUENCE CHARACTERISTICS:

acids (A) LENGTH: 82 amino a. (B) TYPE: amino acid (D) TOPOLOGY: linear

23

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 558:

Leu Leu Lys Val Leu Cys Ile Leu Pro Val Met Lys Val Glu Asn Glu

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Arg Tyr Glu Asn Gly Arg Lys Arg Leu Lys Ala Tyr Leu Arg Asn Thr

Leu Thr Asp Gln Arg Ser Ser Asn Leu Ala Leu Leu Asn Ile Asn Phe 45 Leu Asp Leu Met Val Asp Thr Tyr Ile Lys Leu 55 Ile Lys His Asp Asp

35

Tyr Thr Ser Lys Ser Glu Leu Pro Thr Asp Asn Ser Glu Thr Val Glu 65

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Asn Thr

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(2) INFORMATION FOR SEQ ID NO: 559

(A) LENGTH: 95 amino acids SEQUENCE CHARACTERISTICS: Ξ

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(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 559; 9

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Met Val Leu Ile Leu Leu Asn Leu Leu Leu Gly Gln Phe Ser Cys Met 1 15 15

Ser Pro Ala Ser His His Cys His Pro Leu Pro Thr Glu Met Pro Cys 30 ଌ

Asp Ala Leu Leu Pro Lys Pro Ser Ala Asn Ser 50 55 Ala Glu Gly Pro Ala Ser Leu Arg Cys Asn Lys Tyr Val Ser  $90\,$ Gly His Pro Cys Val Gly Phe Ala Ala Val Leu Val Ala Pro Leu Thr 15 40 45 Cys His Cys Gln Gly Leu Tyr Asn Gln Gln Gln Gln Asn Leu His Ala 65 70 75 80 Tyr Tyr Pro Glu Ser Gly Trp Asp Arg Leu Arg Glu Leu Phe Gly Lys 50 55 60 Met Glu Val Pro Pro Pro Ala Pro Arg Ser Phe Leu Cys Arg Ala Leu 1 15 (2) INFORMATION FOR SEQ ID NO: 561: Val Ala Val Ser Ser Xaa 50 Val Phe Thr Val Ile Gly Asp Ala Pro Gly Ala Val Leu Ser Cys Ala  $20 \hspace{1.5cm} 25$ Met Ile Pro Ala Tyr Ser Lys Asn Arg Ala Tyr Ala Ile Phe Phe Ile 1 15 (2) INFORMATION FOR SEQ ID NO: 560: Glu Val Leu Glu Glu Arg Gln Lys Arg Leu Pro Tyr Val Pro Glu Pro 35 40 45 Ě Ξ (1) SEQUENCE CHARACTERISTICS: SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 561: SEQUENCE DESCRIPTION: SEQ ID NO: 560: (A) LENGTH: 108 amino (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 54 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear Phe Pro Asn Gly Ser 60

Ser Ser Asp Txp Gly Phe Asp Ser His Thr Val Tyr Pro Ser Cys Val
35 40 45

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8 Asp Thr Val Asn Thr Ser Leu Asn Val Tyr Arg Asn Lys Asp Ala Leu 65 70 75

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Ser His Phe Val Ile Ala Gly Ala Val Thr Gly Ser Leu Phe Arg Ile 85 90 95

Asn Val Gly Leu Arg Gly Trp Trp Leu Val Ala Xaa 100 105

(2) INFORMATION FOR SEQ ID NO: 562:

5

(i) SEQUENCE CHARACTERISTICS

Ě (A) LENGTH: 50 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 562:

15

20 Met Asn Trp Gly Leu Ser Ile Trp Leu His Tyr Tyr Glu Lys Lys Lys 1 5 10 Glu Gln Val

Ser Asp Gly Ile Leu Gln Phe Glu Ser Ser Leu Leu Lye Met Arg Arg 35 Phe Leu Val Ile Leu Ala His Val Val Arg Arg Cys Ala 20 25 30

Ala Pro 50

25

(2) INFORMATION FOR SEQ ID NO: 563:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 563: (i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 253 amino acids

35

Met Val Lys Val Cys Asn Asp Ser Asp Arg Trp Ser Leu Ile Ser Leu · 1 15 Asn Ą Ser Gly Lys Asn Val Glu Leu Lys Phe Val Asp Ser Leu 20 25 30

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Arg Gln Phe Glu Phe Ser Val Asp Ser Phe Gln Ile Lys Leu Asp  $$\rm 35$ 

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Leu Leu Leu Phe Tyr Glu Cys Ser Glu Asn Pro Met Thr Glu Thr 50 55

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Phe His Pro Thr Ile Ile Gly Glu Ser Val Tyr Gly Asp Phe Gln Glu 65 70 75 80 Ala Phe Asp His Leu Cys Asn Lys Ile Ile Ala Thr Arg Asn Pro Glu 85 90 95

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Gly Phe Arg Pro Ala Ser Asp Glu Ile Lys Thr Leu Gln Arg Tyr Met Glu Ile Arg Gly Gly Gly Leu Leu Lys Tyr Cys 100 105 Asn Leu Leu Val Arg 110

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Cys Ser Arg Phe Phe Ile Asp Phe Ser Asp Ile Gly Glu Gln Gln Arg 130 Lys Leu Glu Ser Tyr Leu Gln Asn His Phe Val Gly Leu Glu Asp Arg 145 120 115

Lys Tyr Glu Tyr Leu Met Thr Leu His Gly Val Val Asn Glu Ser Thr 175

Met Gly His Glu Arg Arg Gln Thr Leu Asn Leu Ile Thr 180

Val Cys Leu

2

Met Leu Ala Ile Arg Val Leu Ala Asp Gin Asn Val Ile Pro Asn Val 195 2

Asn Phe Ser Asn Tyr Tyr Ile Ala Gln Val Gln Pro Val Phe Thr Cys 235 Ala Asn Val Thr Cys Tyr Gln Pro Ala Pro Tyr Val Ala Asp Ala 210

2

Gln Gln Gln Thr Tyr Ser Thr Trp Leu Pro Cys Asn Xaa 250

22

(2) INFORMATION FOR SEQ ID NO: 564:

SEQUENCE CHARACTERISTICS:
(A) LENGTH: 18 amino acids (B) TYPE: emino acid (D) TOPOLOGY: linear 39

Met Ser Phe Leu Met Trp Leu Met Ser Leu Ala Ile Thr Ser Gln Pro

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 564:

35

Pro Met **\$** 

INPORMATION FOR SEQ ID NO: 565: 3 45

(A) LENGTH: 80 amino acids SEQUENCE CHARACTERISTICS:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 565; (B) TYPE: amino acid (D) TOPOLOGY: linear

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Met Ala Pro Lys Gly Lys Val Gly Thr Arg Gly Lys Lys Gln Ile Phe 1 5 Glu Glu Asn Axg Glu Thr Leu Lys Phe Tyr Leu Axg Ile Ile Leu Gly 20 Ala Asn Ala Ile Tyr Cys Leu Val Thr Leu Val Phe Phe Tyr Ser Ser 40

8

Ala Ser Phe Trp Ala Trp Leu Ala Leu Gly Phe Ser Leu Ala Val Tyr 50 60

Gly Ala Ser Tyr His Ser Met Ser Ser Met Ala Arg Ala Ala Phe Phe 65

9

INFORMATION FOR SEQ ID NO: 566: 3

(A) LENGTH: 73 amino acids SEQUENCE CHARACTERISTICS: TYPE: amino acid

15

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 566: (D) TOPOLOGY: linear

8

His Leu Lys Asp Val Ile Leu Leu Thr Ala Ile Val Gln Val Leu Ser

Cys Phe Ser Leu Tyr Val Trp Ser Phe Trp Leu Leu Ala Pro Gly Arg 20

22

Ala Leu Tyr Leu Leu Trp Val Asn Val Leu Gly Pro Trp Phe Thr Ala

Asp Ser Gly Thr Pro Ala Pro Glu His Asn Glu Lys Arg Gln Arg Arg 50 60

3

Gln Glu Arg Arg Gln Met Lys Arg Leu 65

33

(2) INFORMATION FOR SEQ ID NO: 567:

(A) LENGTH: 263 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear SEQUENCE CHARACTERISTICS: 3

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 567:

Met Asp Cys Pro Ala Leu Pro Pro Gly Trp Lys Lys Glu Glu Val Ile 1 5 45

Gly Leu Ser Ala Gly Lya Ser Asp Val Tyr Tyr Phe Ser 20 Arg Lys Ser S

Ser Gly Lys Lys Phe Arg Ser Lys Pro Gln Leu Ala Arg Tyr Leu 35 40 Lys Met Gly Asn Thr Val Asp Leu Ser Ser Phe Asp Phe Arg Thr Gly 50 60

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Met Pro Ser Lys Leu Gln Lys Asn Lys Gln Arg Leu Arg Asn Asp Pro 65 78

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Leu Arg Gln Ser Gln Gly Lys Ser Pro Leu Leu Trp Lys Arg Thr Leu  $20 \ 25 \ 30$ Oln Gly Val Gly Pro Gly Ser Asn Asp Glu Thr Leu Leu Ser Ala Val  $165\,$ His Pro Ser Asn Lys Val Lys Ser Asp Pro Gln Arg Met Asn Glu Gln 115 120 125 λrg Met Met Arg Pro Phe Tyr Leu Leu Leu Pro Val Leu Cys 1  $^{\circ}$  . 5 Ala Ser Ala Leu His Thr Ser Ser Ala Pro Ile Thr Gly Gln Val Ser 180 180 Asp Val Thr Glu Gln Ile Ile Lys Thr Met Glu Leu Pro Lys Gly Leu 145 150 150 Pro Gln Met Lys Thr Ser Gly Asn 50 55 Leu Phe Gly Leu Thr His Leu 35 Glu Arg Val Gln Gln Val Arg Lys Lys Leu Glu Glu Ala Leu Met Ala 225 230 230 Ę Ala Ala Val Glu Lys Asn Pro Ala Val Trp Leu Asn Thr Ser Gln Pro 195 200 205 (2) INFORMATION FOR SEQ ID NO: 568: ₫ Asp Ile Leu Gln Thr Ala Ser Ile Phe Lys Gln Pro Val Thr Lys Val Thr Asn 100 105 110 Cys Lys Ala Phe Ile Val Thr Asp Glu Asp Ile Arg Lys Gln Glu 210 \$215\$Arg Gln Leu Phe Trp Glu Lys Arg Leu Gln Gly Leu Ser Ala Ser 130 140 Ser Gly Asp Glu Ala Xaa 260 Ě (i) SEQUENCE CHARACTERISTICS: Ser Arg Ala Ala Asp 245 SEQUENCE DESCRIPTION: SEQ ID NO: 568: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 70 amino acids Arg Lys Ser Glu Tyr Ser Lys Tyr Ala 60 Asn Pro Ser Ala Lys Leu Leu Ser 40 45 Thr Glu Glu Met Asp Ile Glu Met 250 255 e Thr Gln Ala 15

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Tyr Asp Thr

Pro Pro Pro Asp Cys His Cys His Ser Phe Arg Ala Glu 20 25 30

Met Pro Val Thr Ser Lys Arg Thr Leu 1

Phe Phe Pro Asp Pro Cys Ser

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 569:

(A) LENGTH: 34 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(2) INFORMATION FOR SEQ ID NO: 569:

(1) SEQUENCE CHARACTERISTICS

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Leu Leu

INFORMATION FOR SEQ ID NO: 570:

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E Ě SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 570: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 104 amino acids

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Met Asn Ser Arg Gly Met Trp Leu Thr Tyr Ala Leu Gly Val Gly Leu 1 15 Leu His Ile Val Leu Leu Ser Ile Pro Phe Phe Ser Val 20 25 l Pro Val Ala 30

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His Ala Val Lys Gly Thr Pro Phe Glu Thr Pro Asp Gln Gly Lys Ser  $50 \ \ \, 55$ 

Trp Thr Leu Thr Asn Ile Ile His Asn Leu Gly Met Tyr Val Phe Leu  $35 \ 40 \ 45$ 

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lle Phe Thr Glu Val Phe His Asn Phe Ser Asn Asn Ser Ile Phe Ser 95 Lys Ala Pro Asn Ser Leu Gly Thr Thr Gly Leu Trp Ser Thr Val Tyr 65 70 75 80

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8 Gly Lys Phe Leu Tyr Glu Val Xaa 100

2 INFORMATION FOR SEQ ID NO: 571:

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(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 132 amino acids
(B) TYPE: amino acid

ξ SEQUENCE DESCRIPTION: SEQ ID NO: 571: (D) TOPOLOGY: linear

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Asn Trp Lys Lys His 1

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Met Trp Leu Thr Tyr Ala Leu Gly Val Gly Leu Leu His Ile Val Leu 1 5 10 15 Leu Ser Ile Pro Phe Phe Ser Val Pro Val Ala Trp Thr Leu Thr Asn 20 25 S

Ile Ile His Asn Leu Gly Met Tyr Val Phe Leu His Ala Val Lys Gly 35

Thr Pro Phe Glu Thr Pro Asp Gln Gly Lys Ala Arg Leu Leu Thr His 50 60

2

Trp Glu Gln Leu Asp Tyr Gly Val Gln Phe Thr Ser Ser Arg Lys Phe 65 Phe Thr lle Ser Pro Ile Ile Leu Tyr Phe Leu Ala Ser Phe Tyr Thr 95

15

Pro Thr His Phe Ile Leu Asn Thr Ala Ser Leu Leu Ser 100 Lys Tyr Asp

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Val Leu Ile Pro Lys Met Pro Gln Leu His Gly Val Arg Ile Phe Gly 120

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Ile Asn Lys Tyr 130

(2) INFORMATION FOR SEQ ID NO: 572:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 572. SEQUENCE CHARACTERISTICS:
(A) LENGTH: 12 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Asn Lys Trp Ile Cys Glu Met His Cys Tyr Leu Val Leu Leu Ser 1 10 15 <del>4</del> Val Cys Ser Pro Ser Ala Leu Arg Arg Val Arg His Thr Leu Ser Arg 20 25

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(2) INFORMATION FOR SEQ ID NO: 573: S (A) LENGTH: 28 amino acids

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 573:

Met Pro Val Leu Ser Leu Leu Cys Thr Leu Ile Val Ser Phe Gln Ser 1 10 15

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Ala Asp Ser Cys Glu Val Phe Leu Asn Cys Ser Leu 25

(2) INFORMATION FOR SEQ ID NO: 574:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 40 amino acids

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(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 574:

Met Lys Val Ser Thr Met Leu Trp Phe Leu Cys Trp Glu Gln Ser His 1 10 15

15

Phe Leu Arg Glu Trp Glu Asp Leu Ser Thr Phe Leu Ile Leu Ile Gln 20 20

Met Glu Cys Gln Tyr Gly Asn Ser 35

20

(2) INFORMATION FOR SEQ ID NO: 575:

25

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 30 amino acids

(B) TYPE: amino acid

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(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQUENCE 575;

Pro Ala Ser Met Gly Leu Pro Leu Met Ala Leu Met Trp Ser Thr Leu 1

Ala Gly Val Asn Phe Ile Leu Ala Leu Pro Leu Leu Xaa Leu 20 30

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(2) INFORMATION FOR SEQ ID NO: 576:

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 29 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 576;

Met Lys Arg Gly Cys Leu Gly Leu Leu Phe Phe Ser Cys Cys Ser Ser 1 1 15

20

Ala Pro Thr Met Leu Leu Cys Asp Tyr Leu Asn Trp Phe 20

(2) INFORMATION FOR SEQ ID NO: 577:

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(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 92 amino acids

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 577: (B) TYPE: amino acid
(D) TOPOLOGY: linear

S Met Lys Leu Leu Gly Ile Ala Leu Leu Ala Tyr Val Ala Ser Val 1 5

Lys Ile Glu Ser Lys Ile Glu Glu Met Val Glu Pro Leu Arg Glu Lys \$35\$

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5 Ile Arg Asp Leu Glu Lys Ser Phe Thr Gln Lys Tyr Pro Pro Val Lys  $50 \ \ \,$ 

(2) INFORMATION FOR SEQ ID NO: 578:

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(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Tyr Ser Arg Ala Asp Glu Val Val Leu Ala Arg Asp Ile Glu Arg Met 50 55 60

Val Glu Ala Arg Leu Ala Arg Arg Val Leu Ala Arg Ser Val Asp Phe 65 70 75

Tyr Asp Arg Leu Gln Asp Ala Gly Ser Arg Trp Pro Glu Leu Tyr Leu  $35 \ 40 \ 45$ 

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Met Leu Arg Leu Leu Leu Leu Val Ala Phe Ala Leu Val Val Val Leu 1 10 15

(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 580:

Phe His Val Leu Leu Ala Pro Ile Thr Ala Leu Phe His Thr His Phe  $20 \ 25 \ 30$ 

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INFORMATION FOR SEQ ID NO: 580:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 110 amino acids (B) TYPE: amino acid

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Lys Phe Asn Lys Lys Lys 65 70

30

SEQUENCE DESCRIPTION: SEQ ID NO: 578:

Met Lys 1

፩ Leu Ile Phe Leu Phe Pro Leu Ser Phe Leu Pro Asn Ile Trp Arg 20 25 30

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4 ρıς Va. Met Val Asn Leu Asn Ile Leu Phe 35

(2) INFORMATION FOR SEQ ID NO: 579:

Leu Ala Gin Glu Cys Pro Pro His Ile Pro Ser Ser Phe Phe Leu Val 1 , 5

Lys Leu Leu Phe Ile Pro Trp Leu Ala Ser Leu Leu Pro Pro Leu Ser  $20 \ \ 30$ 

3

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 70 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 579:

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Thr Phe Thr Ser Asp Phe Tyr Phe Met Glu Phe Gly Ile Glu Val Lys  $_{15}^{\circ}$   $_{15}^{\circ}$   $_{45}^{\circ}$ 

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Trp Gly Asn Phe Val Asn Met Arg Ser Ile Gln Glu Asn Gly Glu Leu  $20 \ \ 30$ 

20 Phe Leu Ser Glu Lys Asp Arg Lys Arg Ile Leu Xaa Asn Arg Arg Arg 65 70 75

Xaa Val Arg Gly Leu Pro Ser Xaa Leu Thr Asn Ser 85 90

E SEQUENCE CHARACTERISTICS (A) LENGTH: 42 amino acids

Phe Ser Leu Val Leu Leu Ile Lys Ile Ile Ser Phe Glu Arg 5 10 15

Val Ser Ser Ala His Val Ser His Leu 85

Arg Asp Tyr Pro Thr Tyr Tyr 90 95

ઝ Thr Ser Leu Cys Val Asp Phe Met Arg Asn Cys Val Arg Cys
100 105 110

6 (2) INFORMATION FOR SEQ ID NO: 581:

(i) SEQUENCE CHARACTERISTICS

(A) LENCTH: 30 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 581:

3

Met Phe Lys Leu Glu Glu Cys Gly Lys Thr Thr Phe Leu Leu Ser Met
1 10 15

Ala Leu Tyr Phe Trp Trp Ile Val Gln Thr Thr Lys Gly Cys
25 30

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3 INFORMATION FOR SEQ ID NO: 582:

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(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 71 amino acids

(B) TYPE: amino acid

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Leu Gln Gln Cys Arg Gln His Gln Val Leu Gln Glu Lys Asn Thr Lys 50 60

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 582: (D) TOPOLOGY: linear

Met Glu Ser Asp Ala Leu Leu Leu Thr Ile Phe Trp Ile Ile Ala Arg

S

Ser Ser Val Arg Ser Val Gly Lys Ser Ser Gln Arg Ser Phe Thr Thr 30

Ile Thr Gln Leu Arg Ser Thr His Thr Gly Pro Ser Arg Arg Ser Tyr 2 Leu lle Trp Trp Asn Gly Gly Pro Lys Arg Thr lle Ser Tyr Val Ser 50

Arg Arg Phe Arg Ser Phe Arg 65 70

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(2) INFORMATION FOR SEQ ID NO: 583:

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SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 583: (D) TOPOLOGY: linear

val Gly Leu Phe Gln Pro Lys Thr Phe Gln Val Pro Val Thr Asp Leu

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lyr ile Phe ile Lys ile Tyr Ser Glu ile Gly Pro ile Met His Val

Pro Gly Tyr Ser Gln Ser Pro Ser Thr Pro Pro Trp Thr Leu cys 35

(2) INFORMATION FOR SEQ ID NO: 584; 6

SEQUENCE CHARACTERISTICS:

TYPE: amino acid

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SEQUENCE DESCRIPTION: SEQ ID NO: 584:

Met Trp Phe Gly Ser Asp Arg Ser Asp Leu Arg Ile Gly Thr Ala Phe 1 5 10 S Leu Phe Asp Leu Val Cys Asp Leu Cys Ile His Ala Trp Lys Pro Pro

Gly Leu Val Arg Phe Ser Phe

55

(2) INFORMATION FOR SEQ ID NO: 585:

S

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 19 amino acids

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 585: (D) TOPOLOGY: linear

Leu Asn Thr Ala Ser Leu Asn Leu Pro Trp Lys Val Gln Leu Phe 10 10 15

Ala His Ala 2

(2) INFORMATION FOR SEQ ID NO: 586: 2 (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 586:

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Met Ser Ala Cys Leu Leu Leu Phe Leu Ala Phe Ser Trp Lys Arg Lys

Gly Leu Trp Ser Gly Pro Gly

22

8

(2) INFORMATION FOR SEQ ID NO: 587:

(A) LENGTH: 69 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

33

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 587:

Leu Pro Pro Phe Ser Leu Val Tyr Thr His Phe Leu Val Ala Ser  $$\rm 10$ Met

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Leu Leu Pro Val Ile Leu Ala Val Phe Pro Asp Ser Ala Gln Ile Val

Pro Leu Lys Pro Ile Pro Arg Pro Gln Pro Glu Val Ile Phe Pro 35 45

Ser Ser Glu Leu Leu Glu Gln Leu Leu Ser Val Gln Phe Val Trp Gln

Ala His Thr Val Ala

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(2) INFORMATION FOR SEQ ID NO: 588:

55

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 77 amino acids (B) TYPE: amino acid

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Thr Leu Ser Thr Phe Lys Glu Ser Trp Lys Xaa 145 150 155

Gln Leu Leu Ala,Gln Thr Lys Lys Val Val Arg Pro Thr Arg Lys Lys 130 140

Ile Pro Ala Ile Val Cys Leu Gly Phe Leu Leu Leu Leu Asn Val Gly
115 120 125

Gly Ala Ile Phe Thr Leu Ala Ala Leu Lys Glu Ser Leu Ser Thr Cys 100 105

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2 INFORMATION FOR SEQ ID NO: 590:

(1) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 amino acids

(B) TYPE: amino acid
(D) TOPOLOGY: linear

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Ě SEQUENCE DESCRIPTION: SEQ ID NO: 590:

Net Pro Glu Thr Arg Leu Gly His Arg Gln Gln Phe Ala Val Phe His 1 10 15

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Ē Xaa Pro Val Pro Pro Cys Gly

(2) INFORMATION FOR SEQ ID NO: 591:

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20 (i) SEQUENCE CHARACTERISTICS

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(2) INFORMATION FOR SEQ ID NO: 589:

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SEQUENCE CHARACTERISTICS (A) LENGTH: 155 amino e (B) TYPE: amino acid (D) TOPOLOGY: linear

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Met Ala Leu Leu Leu Sar Val Leu Arg Val Leu Leu Gly Gly Phe Phe
1 10 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 589:

Ala Leu Val Gly Leu Ala Lys Leu Ser Glu Glu Ile Ser Ala Pro Val 20

5

Asn Val Gly Gln Leu Leu Ala Gln Thr Lys Lys Val Val Arg Pro Thr  $50 \ \,$ 

Arg Lys Lys Thr Leu Ser Thr Phe Lys Glu Ser Trp Lys 65

Ser Thr Cys Ile Pro Ala Ile Val Cys Leu Gly Phe Leu Leu Leu Leu 35

Leu Met Met Gly Ala Ile Phe Thr Leu Ala Ala Leu Lys Glu Ser Leu 20 30

5

S

Met Gly Pro Pro Met Leu Gln Glu Ile Ser Asn Leu Phe Leu Ile Leu 1 15

(D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 588:

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 591: (A) LENGTH: 38 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Met Lou Thr Phe Leu Phe Ser Ala Cys Ala Thr Cys Leu Gly Lys Leu 1 15

23

Gly Pro Pro Leu Leu Ser 35 Ala Ser Pro Leu Ala Pro Val Gly Pro Gln Gln Arg Gly Xaa Pro Pro 20 25 30

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(2) INFORMATION FOR SEQ ID NO: 592:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 592: (1) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 69 amino acids

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Ser Glu

Arg Met Asn Ala Leu 35

Phe Val Gln Phe Ala Glu Val Phe Pro 40

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Pro Met Leu Gln Glu Ile Ser Asn Leu 85

Phe Leu Ile Leu Leu Met Met 90 95

Val Gly Phe Leu Glu Leu Leu Ala Gly Leu Leu Leu Val Met Gly Pro 65 70 75

Leu Lys Val Phe Gly Tyr Gln Pro Asp Pro Leu Asn Tyr Gln Ile Ala 50

45 Met Asp Pro Phe His Tyr Asp Tyr Gln Thr Leu Arg Ile Gly Gly Leu 10 15

Val Phe Ala Val Val Leu Phe Ser Val Gly Ile Leu Leu Ile Leu
20 25 30

50 Arg Arg Cys Lys Cys Ser Phe Asn Gln Lys Pro Arg Ala Pro Gly Asp 35

Glu Glu Ala Gln Val Glu Asn Leu Ile Thr Ala Asn Ala Thr Glu Pro 50 55 60

Gln Lys Ala Glu Asn 65

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			-\$70 A 1 869	
	(2) INFORMATION FOR SEQ ID NO: 593:		275 280 285	
ν,	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 308 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 593:	S	Glu Ile Leu Gln Ile His Thr Lys Asn Met Lys Leu Ala Asp Asp Val 290 295 Asp Leu Glu Gln 305	
0	Asn Leu Arg Val Arg Leu Gly Asp Val Ile Ser Ile Gln Pro Cys Pro 1 15 15		TO ON THE OWN THAT AND THE TAXABLE INC.	
	Asp Val Lys Tyr Gly Lys Arg lle His Val Leu Pro Ile Asp Asp Thr 20 30		(2) INFURMATION FOR SECTION SONS: (1) SEQUENCE CHARACTERISTICS:	
2	Val Glu Gly Ile Thr Gly Asn Leu Phe Glu Val Tyr Leu Lys Pro Tyr 35 46		(A) LENGTH: 22 amino acids (B) TYPE: amino acid (D) TOPLOGY: linear	
_	Phe Leu Glu Ala Tyr Arg Pro Ile Arg Lys Gly Asp Ile Phe Leu Val 50		Met Gln	
>	Arg Gly Gly Met Arg Ala Val Glu Phe Lys Val Val Glu Thr Asp Pro 65 75 80	07	Leu Leu Val I	
S	Ser Pro Tyr Cys Ile Val Ala Pro Asp Thr Val Ile His Cys Glu Gly 85	25		•
	Glu Pro Ile Lys Arg Glu Asp Glu Glu Glu Ser Leu Asn Glu Val Gly 100		(2) INFORMATION FOR SEQ ID NO: 595:	
0	Tyr Asp Asp Ile Gly Gly Cys Arg Lys Gln Leu Ala Gln Ile Lys Glu 115	30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 amino acids (B) TYPE: amino acid	,
U	Met Val Glu Leu Pro Leu Arg Hís Pro Ala Leu Phe Lys Ala Ile Gly 130	•	(X1) SEQUENCE DESCRIPTION: SEQ ID NO:,595:	
_	Val Lys Pro Pro Arg Gly Ile Leu Leu Tyr Gly Pro Pro Gly Thr Gly 145	G.	Met Phe Pro Lys Phe Cys Pro Ile Leu Ser Leu Val Asp Phe Ile Ser 1 5	
_	Lys Thr Leu Ile Ala Arg Ala Val Ala Asn Glu Thr Gly. Ala Phe 175	40	His Arg Asp Lys Pro Glu Thr Glu	
	Phe Leu Ile Asn Gly Pro Glu Ile Met Ser Lys Leu Ala Gly Glu Ser 180		(2) INPORMATION FOR SEQ ID NO: 596:	
10	Glu Ser Asn Leu Arg Lys Ala Phe Glu Glu Ala Glu Lys Asn Ala Pro 200	45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 24 amino acids	
_	Ala ile ile Phe ile Asp Glu Leu Asp Ala ile Ala Pro Lys Arg Glu 210	O <sub>2</sub>	(B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIFIION: SEQ ID NO: 596:	
	Lys Thr His Gly Glu Val Glu Arg Arg Ile Val Ser Gln Leu Leu Thr 225 240		Met Leu Ile Glu Cys Ala Trp Gln Leu Met Phe Leu Leu Leu Lys Val 1 5	· .
	Leu Met Asp Gly Leu Lys Gln Arg Ala His Val Ile Val Met Ala Ala 245	55	Glu Gln Leu Gly Ile Leu Asp Lys 20	
	Thr Asn Arg Pro Asn Ser Ile Asp Pro Ala Leu Arg Arg Phe Gly Arg 260			
_	Phe Asp Arg Glu Val Asp Ile Gly Ile Pro Asp Ala Thr Gly Arg Leu	09	(z) infokkajion fok seq id no: 597:	

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1
                                                                                    Met Xaa Gly Leu Ser Leu Ile Leu Thr Val Thr Leu Leu Ala Val Ser
1 10
                                                                                                                                                                                                                                                                                                                                                                       Met Phe Leu Val Trp Phe Phe Trp Gly Leu Ile Ser Ala Leu Ser Asn
1 10 15
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                                  Asp Ser Ala Ala Thr Cys Ile Val Ala Lys Gly
20 , 25
                                                                                                                                                                                                                                                                                                                     Val His Thr Pro Ser Arg Leu Pro Ala
20 25
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                                                                                                                                      SEQUENCE DESCRIPTION: SEQ ID NO: 600:
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              (B) TYPE: amino acid
(D) TOPOLOGY: linear
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                (D) TOPOLOGY: linear
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                                                                                                                                                         (D) TOPOLOGY: linear
                                                                                                                                                                           (B) TYPE: amino acid
                                                                                                                                                                                           (A) LENGTH: 27 amino acids
                                                                                                                                                                                                                                                                                                                                                                                                                                             (D) TOPOLOGY: linear
                                                                                                                                                                                                                                                                                                                                                                                                                                                                             (A) LENGTH: 25 amino acids
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   (A) LENGTH: 8 amino acids
                                                                                                                                                                                                                                                                                                                                                                                                                                                              (B) TYPE: amino acid
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30 23 20 2 5 ઝ S Met Trp Thr Arg Ser Ser Arg Cys Leu Leu Leu Cys Ile Pro Gly Xaa 1 15 Ser Arg Arg Arg Ala Gly Ser Gly Met Lys Pro Arg Ser Trp Ser 25 Thr Gln Ser Arg Thr Leu Ser Ala Thr Ala Ser Pro Ala 50 55 Ser Thr Ser Gly Leu Ser Ala Phe Met Gly Leu Gly Thr 20 25 Met Ary Glu Thr Ser Ile Ary Val Leu Leu Met Leu Pro Ala Leu Glu 1 15 Ala Trp (2) INFORMATION FOR SEQ ID NO: 602: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 601: Arg Pro Ser Gly Gly Thr Gly Thr Ser Ser Ser Gln Ser Ser 35 £ (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 602: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 29 amino acids (D) TOPOLOGY: linear (B) TYPE: amino acid

(2) INFORMATION FOR SEQ ID NO: 603:

Met Pro Pro Lys Gln Glu Leu Gly Ser Gly Val Gly Glu Leu Ala Lys Leu Ile Arg Trp Glu Asp Gly Leu Leu Leu Glu Gly Leu Leu Leu Val
35 40 45 Asm Ser Lys Arg Gln His Trp Asm His Arg Trp Lys Lys Tyr Leu Lys 20 25 30 Ξ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 603: SEQUENCE CHARACTERISTICS: (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 69 amino acids

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Glu His Cys Ala Thr Met Ala Trp Asp Cys Leu Met Arg Leu Glu 50  $$5^\circ$$ 

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(2) INFORMATION FOR SEQ ID NO: 604:

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Leu Leu Lys Arg Leu 65

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INFORMATION FOR SEQ ID NO: 601:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 61 amino acids

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	Arg Val Pro Arg Asp Ile Pro Ala Met Leu Pro Ala Ala Arg Leu Pro 100	
<b>.</b>	Thr Thr Val Leu Asn Ala Thr Ala Lys Ala Val Ala Val Thr Leu Gin 125	
	Ser His · 130	<del>.</del>
10		
	(2) INFORMATION FOR SEQ ID NO: 607:	
15	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 23 anino acids (B) TYPE: anino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 607:	
20	Met Leu Val Ile Phe Leu Phe Thr Ser Leu Leu Lys Ile Pro Ser Ser 1 $^{\rm 1}$	
25	Val Pro Gly Leu Ile Asn Val 20	
	(2) INFORMATION FOR SEQ ID NO: 608:	
. 30	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 6 amino acids (B) TYPE: amino acid (D) TOPPLICSY: linear	
35	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 608:	· · · · · · · · · · · · · · · · · · ·
	Glu Leu Asp Tyr I.e Leu 1	
40	(2) INFORMATION FOR SEQ ID NO: 609:	
45	(i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 232 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (xi) SEQUENCE DESCRIFTION: SEQ ID NO: 609:	·
50	Met Ala Pro Pro Gly Trp Gln Xaa Xaa Xaa Xaa Trp Leu Ala Cys 1 15	
	Pro Asp Arg Gly Glu Leu Ser Ser Arg Ser Pro Pro Cys Arg Leu Ala 20	·
55	Arg Trp Ala Glu Gly Asp Arg Glu Thr Arg Thr Cys Leu Leu Glu Leu 15	-
09	Ser Ala Gin Ser Trp Gly Gly Arg Phe Arg Arg Ser Ser Ala Val Ser 50 60	

Met Asn Leu His Gln Arg Arg Leu Leu Leu Ile Gly His Leu Met Thr

22

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 605:

(i) SDQUDNCE CHARACTERISTICS:
(A) LENGTH: 35 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 605:

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Leu Val Lys Ala Ser Lys Ser Phe Ser Phe Thr Glu Ile Thr Ser Ser 20 25

Arg Lys Lys 35

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Leu Leu Gly Tyr Gly Leu Phe Gly His Cys Ile Val Leu Phe Ile Thr  $_{\rm 1}$   $_{\rm 1}$ 

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(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 130 amino acids
(B) TYPE: amino acid
(D) TYPE: amino acid
(D) TOPLOGY: linear
(Xi) SEQUENCE DESCRIPTION: SEQ ID NO: 606:

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(2) INFORMATION FOR SEQ ID NO: 606:

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Tyr Asn Ile His Leu His Ala Leu Phe Tyr Leu Phe Try Leu Leu Val 20 Gly Gly Leu Ser Thr Leu Arg Met Val Ala Val Leu Val Ser Arg Thr 35 40

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His Met Leu Phe Leu Tyr Leu His Phe Ala Tyr His Lys Val Xaa 65 70 80

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Val Gly Pro Thr Gln Arg Leu Leu Cys Gly Thr Leu Ala Ala Leu 50 Glu Gly Ile Leu Asp Thr Leu Glu Gly Pro Asn Ile Pro Pro Ile Gln 85 85

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Lyg lle Val Tyr lle Leu Gly Asn Pro Leu Lys Phe Asn Ser Arg Val 1 5 10 15

Ile His His Leu Val Leu Leu Gln 20

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(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 24 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: 11near
(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 604:

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Trp Leu Asp . 225 Leu Glu Thr Leu Asp Gly Gly Gln Glu Asp Gly Ser Glu Ala Asp Arg 145 150 150 Pro Pro Pro Pro Arg Met Thr Ser Pro Ser Ser Ser Pro Val Phe Arg 130 140 Thr Leu Asp Ile Ser Ile Cys Thr Glu Val Leu Ala Gly Thr Glu Gln 115 120 125 Gly Leu Pro Val Arg Pro Thr Pro Thr Arg Thr Gly Glu Glu Asp Arg 100 100 Arg Ser Ser Gly 11e Pro Ala Ala Ala Thr Pro Trp Pro Gln Pro Ala 95 Ala Gly Ser Pro Ser Arg Leu His Phe Leu Pro Gln Pro Leu Leu Leu 65 70 75 Thr Glu Ile Gly Ser Ser Met Arg Ser Pro Gly Val Ser Pro Arg Ile 210 225 Gln Gly Glu Asp Arg Lys Phe Ala Pro Ser Asp Lys Ser Gln Pro Pro 180 185 Gly Lys Leu Met Val Leu Leu Leu Leu Ala Tyr Val Leu Leu Thr Tyr Ile Leu 1 5 Thr Thr Glu Arg Glu Gln Val Pro Val Ser Arg Ile Gln Thr Asp Leu 195 200 205 Arg His Leu Leu Asn Met Leu Ile Ala Leu Met Xaa Arg Asp Arg Gln Gln Cys 20 30 (2) INFORMATION FOR SEQ ID NO: 610: XX (1) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 610: Phe Gln Ser Thr Xaa 230 g S (D) TOPOLOGY: linear (A) LENGTH: 34 amino acids
(B) TYPE: amino acid Phe Gly Ser Gly Leu Pro Pro Met Glu Ser Gln Phe 165 170 175

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15 5 \$ 6 ઝ 30 25 20 Gly Pro Cys Trp Pro Ser Thr Gly Asn Cys Cys Leu Cys
20 25 Met Val Trp Val Leu Trp Ser Ala Pro Ser Leu Ala Pro Pro Trp Val
1 15 Gly Lys Lys Asn Gln Leu Leu Val Ile (2) INFORMATION FOR SEQ ID NO: 612: (2) INFORMATION FOR SEQ ID NO: 614: (2) INFORMATION FOR SEQ ID NO: 613: ξ (i) SEQUENCE CHARACTERISTICS: Ĕ (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 613: (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 612: SEQUENCE DESCRIPTION: SEQ ID NO: 614: (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 9 amino acids (D) TOPOLOGY: linear (B) TYPE: amino acid (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 30 amino acids (A) LENGTH: 29 amino acids

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Pro Thr Ser His Pro 20

Met Val Phe Glu Gly Phe Ser Ser Ala Phe Cys Leu Ser Ser Thr Ala 1 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 611:

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INFORMATION FOR SEQ ID NO: 615:

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Cys Cys Gln Pro Cys Arg Pro Thr Ser Ser Ala Pro Met Arg

Met Ala Lys Arg Ser Pro Gly Gly Cys Gly Ser Gly Leu Ile Leu Leu 1 15

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SEQUENCE CHARACTERISTICS:

(A) LENGTH: 21 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

INFORMATION FOR SEQ ID NO: 611:

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(i) SEQUENCE CHARACTERISTICS:

(A) LEWSTH: 113 amino acids (B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 615: (D) TOPOLOGY: linear

Ile Thr Ile Ala Ile Gln Met Ile Cys Leu Val Asn Xaa Glu Leu Tyr

Pro Thr Phe Val Arg Asn Xaa Gly Val Met Val Cys Ser Ser Leu Cys 20

Asp Ile Gly Gly Ile Ile Thr Pro Phe Ile Val Phe Arg Leu Arg Glu 35 2

ž z Leu Gly Leu lle Leu Phe Ala Val 55 60 Val Trp Gln Ala Leu Pro 50 15

Leu Leu Leu Pro Glu Thr Lys Gly Val Ala Leu 70 75 80 Lys Asp Ala Glu Asn Leu Gly Arg Lys Ala Lys 85 95 95 Ala Ala Gly Val Thr Pro Glu Thr Met 65 2

Lys Glu Asn Thr Ile Tyr Leu Lys Val Gln Thr Ser Glu Pro Ser Gly 100

22

(2) INFORMATION FOR SEQ ID NO: 616: 39

(i) SEQUENCE CHARACTERISTICS:

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 616:

Met Lys Asp Ala Glu Asn Leu Gly Arg Lys Ala Lys Pro Lys Glu 5 10 돮

Asn Thr

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(2) INFORMATION FOR SEQ ID NO: 617: 45

(A) LENGTH: 21 amino acids SEQUENCE CHARACTERISTICS: 3

(B) TYPE: amino acid

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SEQUENCE DESCRIPTION: SEQ ID NO: 617: (D) TOPOLOGY: linear (xi

Pro Arg Val Arg Asn Ser Pro Glu Asp Leu Gly Leu Ser Leu Thr Gly 1 5 55

Asp Ser Cys Lys Leu 20

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(2) INFORMATION FOR SEQ ID NO: 618:

(A) LENGTH: 52 amino acids (i) SEQUENCE CHARACTERISTICS:

(B) TYPE: amino acid

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 618:

Gin Ala Asp Asp Leu Gin Ala Thr Val Ala Ala Leu Cys Val Leu Arg

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Pro Gly Gly Gly Gly Fro Trp Ala Gly Ser Trp Leu Ser Pro Lys Thr  $20\ 25\$  Ala Met Gly Gly Asp Leu Val Leu Gly Leu Gly Ala Leu Arg Arg Arg 35

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Lys Arg Leu Leu 50

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(2) INFORMATION FOR SEQ ID NO: 619:

25

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 232 amino ac (B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 619:

3

Glu Gln Glu Lys Ser Leu Ala Gly Trp Ala Leu Val Leu Ala Xaa Xaa 1 1 10 15

Gly ile Gly Leu Met Val Leu His Ala Glu Met Leu Trp Phe Gly Gly 30 25

35

Cys Ser Ala Val Asn Ala Thr Gly His Leu Ser Asp Thr Leu Trp Leu lle Pro lle Thr Phe Leu Thr lle Gly Tyr Gly Asp Val Val Pro Gly 50 60

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The Met Trp Gly Lys Ile Val Cys Leu Cys The Gly Val Met Gly Val 65 5

Cys Cys Thr Ala Leu Leu Val Ala Val Val Ala Arg Lys Leu Glu Phe  $85\ \,$  95

Asn Lys Ala Glu Lys His Val His Asn Phe Met Met Asp Ile Gln Tyr 100 S

Thr Lys Glu Met Lys Glu Ser Ala Ala Arg Val Leu Gln Glu Ala Trp 115 Met Phe Tyr Lys His Thr Arg Arg Lys Glu Ser His Ala Ala Arg Xaa 130

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His Gin Arg Xea Leu Leu Ala Ala Ile Asn Ala Phe Arg Gin Val Arg 145 8

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Ser Lys Met His Met Ile Leu Tyr Asp Leu Gln Gln Asn Leu Ser Ser 180 185 Leu Lys His Arg Lys Leu Arg Glu Gln Val Asn Ser Met Val Asp Ile 165 170 175 Asp Ala Leu Thr Glu Leu Leu Ser Thr Ala Leu Gly Pro Arg Gln Leu 210 215 Ser His Arg Ala Leu Glu Lys Gln Ile Asp Thr Leu Ala Gly Lys Leu 195 200 205 Thr Arg Gly Gly Phe Arg Gly Cys Thr Val Trp Leu Thr Gly Leu Ser  $20\ 25\$ Pro Glu Pro Ser Gln Gln Ser Lys 230 His Lys Ala Lys Ser His Pro Glu Val 50 55 Cys Asn Ile Cys Gly Lys Lys Phe Glu Lys Lys Asp Ser Val Val Ala 35 . 40  $\phantom{-}45$ Asn Tup His Met Lys Lys His Asp Ala Asp Ser Phe Tyr Gln Phe Ser 25  $30\,$ Leu Gln Cys Glu Ile Cys Gly Phe Thr Cys Arg Gln Lys Ala Ser Leu 1 15 Gly Ala Gly Lys 35 Tyr Gin Ala His His Val Ser Arg Asn Lys Arg Gly Gin Val Val Gly
1 15 2 3 INFORMATION FOR SEQ ID NO: 620: INFORMATION FOR SEQ ID NO: 621: £ (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 620: (i) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 621: SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 57 amino acids (A) LENGTH: 36 amino acids

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\$ 30 50 6 35 25 20 5 7 Ala Gly Pro 50 Ile Thr Ser Thr Asp Ile Leu Gly Thr Asn Pro Glu Ser Leu Thr Gln
1 19 Val Gly Ser Gly Gly Gly Thr Glu Gly Leu Val Met Asn Ser
20 25 30 Cys Ser Gly Thr Glu Arg Val Ser Leu Met Ala Asp Gly Lys Ile Phe 1 15 Asp Ile Leu Gly Ala Thr Thr Glu Val Leu Ile Glu Asp Ser Asp Ser 35 Lys Ser Tyr Asn Ser Thr Ser Gly Glu Cys Leu Leu Leu Glu Ala Glu Gly Met Ser 1 15 . (2) INFORMATION FOR SEQ ID NO: 623: Pro Ser Asp (2) INFORMATION FOR SEQ ID NO: 624: (i) SEQUENCE CHARACTERISTICS: (i) SEQUENCE CHARACTERISTICS: ξ (i) SEQUENCE CHARACTERISTICS (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 624: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 622: SEQUENCE DESCRIPTION: SEQ ID NO: 623: (D) TOPOLOGY: linear (B) TYPE: amino acid (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 51 amino acids (D) TOPOLOGY: linear (A) LENGTH: 19 amino acids (A) LENGTH: 19 amino acids

(i) SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (A) LENGTH: 60 amino acids

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(2) INFORMATION FOR SEQ ID NO: 625:

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 625: (D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 622:

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His Pro Arg Tyr Leu Gln His His Ile Lys Tyr Gln His Leu Leu Lyr  $20\ 20$ Lys Lys Tyr Val Cys Pro His Pro Ser Cys Gly Arg Leu Phe Arg Leu 45 Ile Gln Tyr Val Arg Cys Glu Met Glu Gly Cys Gly Thr Val Leu Ala 1 5 10 Gln Lys Gln Leu Leu Arg His Ala Lys His His Thr 50 50 60

(2) INFORMATION FOR SEQ ID NO: 626:

15

(A) LENGTH: 31 amino acida SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (D) TOPOLOGY: linear 2

Asp Gin Arg Asp Tyr 11e Cys Glu Tyr Cys Ala Arg Ala Phe Lys Ser 1 5 15 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 626; 25

Ser His Asn Leu Ala Val His Arg Met Ile His Thr Gly Glu Lys 20 30

(2) INFORMATION FOR SEQ ID NO: 627:

3

(A) LENGTH: 25 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid 3 35

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 627;

Arg Ser Ser Arg Ser Lys Thr Gly Ser Leu Gln Leu lle Cys Lys Ser 1 5 <del></del>

Glu Pro Asn Thr Asp Gln Leu Asp Tyr 20

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(2) INFORMATION FOR SEQ ID NO: 628:

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(A) LENGTH: 183 amino acids (B) TYPE: amino acid 3

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 628:

Leu Gln Cys Glu Ile Cys Gly Phe Thr Cys Arg Gln Lys Ala Ser Leu

55

Asn Trp His Met Lys Lys His Asp Ala Asp Ser Phe Tyr Gin Phe Ser 20 8

Cys Asn Ile Cys Gly Lys Lys Phe Glu Lys Lys Asp Ser Val Val Ala 35

Asp Ile His Lys Ala Lys Ser His Pro Glu Val Xaa Ile Thr Ser Thr

Lou Gly Thir Asn Pro Glu Ser Leu Thir Gln Pro Ser Asp Xea Asn Ser 65 70 70

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Lys Ser 95 Thr Ser Gly Glu Cys Leu Leu Leu Glu Ala Glu Gly Met Ser

Tyr Xaa Cys Ser Gly Thr Glu Arg Val Ser Leu Met Ala Asp Gly Lys 100 lie Phe Val Gly Ser Gly Ser Gly Gly Thr Gly Gly Leu Val Met 115

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Ser Asp Ile Leu Gly Ala Thr Thr Glu Val Leu Ile Glu Asp Ser 130 Asn ೫

Arg Ala Phe Lys Ser Ser His Asn Leu Ala Val His Arg Met 11e His 175 Asp Ser Ala Gly Pro Xea Gln Arg Asp Tyr Ile Cys Glu Tyr Cys Ala 145 23

Thr Gly Glu Lys His Tyr Xea 180

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(2) INFORMATION FOR SEQ ID NO: 629:

35

(A) LENGIH: 60 amino acids SEQUENCE CHARACTERISTICS: (B) TYPE: amino acid (x1) SEQUENCE DESCRIPTION: SEQ ID NO: 629:

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Gin Tyr Val Arg Cys Glu Met Glu Gly Cys Gly Thr Val Leu Ala His  $_{\rm 1}$ Pro Arg Tyr Leu Gln His His Ile Lys Tyr Gln His Leu Leu Lys Lys 2045

Lys Tyr Val Cys Pro His Pro Ser Cys Gly Arg Leu Phe Arg Leu Gln 35 Lys Gin Leu Leu Arg His Ala Lys His His Thr Asp 50 55 S

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(2) INFORMATION FOR SEQ ID NO: 630:

SEQUENCE CHARACTERISTICS Э

(A) LENGTH: 27 amino acids (B) TYPE: amino acid

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ၓ 25 20 5 45 6 35 15 S 50 S Asp Leu Val His Arg Glu Glu Ala Ser Cys Tyr Cys Gln Ala Glu Pro Ser Gly
1 15 Lys Arg Trp Ala Gly Leu 20 Arg Pro Ala Leu Arg Gln Ala Gly Gly Gly Thr Arg Glu Pro Arg Gln
1 15 (2) INFORMATION FOR SEQ ID NO: 634: Ser Leu Phe Leu Leu Val Val Leu Tyr His Tyr Val Ala Val Asn Asn 20 25 30 Met Ile Thr Asp Val Gln Leu Ala Ile Phe Ala Asn Met Leu Gly Val 1 15 Ala Val Asn Phe Arg Pro Gln Arg Ser Gln Ser Met 1 10 (2) INFORMATION FOR SEQ ID NO: 635: (2) INFORMATION FOR SEQ ID NO: 636: (1) SEQUENCE CHARACTERISTICS: (i) SEQUENCE CHARACTERISTICS: ž (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 635: (i) SEQUENCE CHARACTERISTICS: Ĕ  $\tilde{\mathbf{x}}$ (i) SEQUENCE CHARACTERISTICS: SEQUENCE DESCRIPTION: SEQ ID NO: 633: SEQUENCE DESCRIPTION: SEQ ID NO: 634: SEQUENCE DESCRIPTION: SEQ ID NO: 636: (A) LENGTH: 18 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear (B) TYPE: amino acid (D) TOPOLOGY: linear (A) LENGTH: 22 amino acids (B) TYPE: amino acid
(D) TOPOLOGY: linear (A) LENGTH: 37 amino acids (D) TOPOLOGY: linear (B) TYPE: amino acid (A) LENGTH: 12 amino acids

Glu Met Phe Asp Ser Leu Ser Tyr Phe Lys Gly Ser Ser Leu Leu Leu 1 5 Leu Tyr Leu His Asn His Ser Tyr Ala Ser Ile Gln Ser Asp Asp Leu 35 40 Met Leu Lys Thr Tyr Leu Ser Glu Asp Val Phe Gln His Ala Val Val 20 25

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SEQUENCE DESCRIPTION: SEQ ID NO: 631:

(A) LENGTH: 110 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

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(2) INFORMATION FOR SEQ ID NO: 631:

(1) SEQUENCE CHARACTERISTICS:

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Pro Phe Lys Asp Asp Pro Arg Asp Glu Thr Tyr Lys Pro His Leu Glu 1 15

(x1) SEQUENCE DESCRIPTION: SEQ ID NO: 630:

(D) TOPOLOGY: linear

Arg Glu Thr Pro Lys Pro Arg Arg Lys Ser Gly
20 25

Met Met Lys Thr Trp Thr Leu Gln Lys Gly Phe Pro Leu Val Thr Val 65 70 75

Trp Asp Ser Phe Asn Glu Val Thr Asn Gln Thr Leu Asp Val Lys Arg 50

30

Gln Lys Lys Gly Lys Glu Leu Phe Ile Gln Gln Glu Arg Phe Phe Leu 95

35

Asn

Met Lys Pro Glu Ile Gln Pro Ser Asp Thr Arg Tyr Met 100 105

6 2 INFORMATION FOR SEQ ID NO: 632:

50 3 (1) SEQUENCE CHARACTERISTICS: (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 632: (A) LENGTH: 24 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

Leu Glu Lys Val Ala Ser Val Gly Asn Ser Arg Pro Thr Gly Gln Gln 15

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Leu Glu Ser Leu Gly Leu Leu Ala 20

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(2) INFORMATION FOR SEQ ID NO: 633:

Pro Lys Lys Gln Glu 35

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INFORMATION FOR SEQ ID NO: 637: 3

SEQUENCE CHARACTERISTICS:
(A) LENOTH: 342 amino acids
(B) TYPE: amino acid

(D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 637: Œ

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Phe Leu Gln Asp Leu Ala Arg 10 Glu Glu Met Ala Asp Ser Val Lys Thr

Gly lie Lys Asp Ser lie Trp Gly lie Cys Thr lie Ser Lys Leu Asp \$20\$

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Ala Arg Ile Gin Gin Lys Arg Giu Giu Gin Arg Arg Arg Ala Ser 15

Val Leu Ala Gin Arg Arg Ala Gin Ser Ile Giu Arg Lys Gin Glu 50 60 Ser 2

lle Val Sar Arg Ile Phe Gln Cys Cys Ala Trp Asn 70 70 80 Glu Pro Arg Ser 65

Phe Ser Leu Leu Leu Phe Tyr Arg Val Phe Ile ਸੂਹ 85 Gly Gly Val Phe

25

ŝ Pro Ser Asp Pro Val Leu Gln Ser Val Thr Ala Arg Ile Ile Gly 100 3

Thr Ser Ile Phe Trp Leu Glu Phe Phe Leu Gly Asp Val Trp Ser НİВ

Leu Pro Leu Phe Val Leu Ser Lys Val Val Asn 135 Val Ser Ala Leu Trp 35

Ala Ile Trp Phe Gln Asp Ile Ala Asp Leu Ala Phe Glu Val Ser Gly 145

Pro Phe Pro Ser Val Ser Lys lle lle Ala Asp Met 165 Arg Lys Pro His <del>4</del>

Leu Phe Asn Leu Leu Gan Ala Leu Phe Leu Ile Gln Gly Met Phe 180 45

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Ser Leu

Ser Leu Tyr Cys Phe Glu Tyr Arg Trp Phe 215 Leu Val Leu Phe Pro Ile His Leu Val Gly Gln 195 Met Ser Leu Leu Tyr 210 HÌB ႙

Ser Val

Leu Thr Ala Asn Lys Gly Ile Glu Met His Gln Arg Leu Ser Asn Ile Glu Arg Asn 235 몺 Leu Ala Trp Pro Tyr Tyr Phe Gly Phe Gly Leu Pro 245 250

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Met Gln ser ser Tyr 11e 11e ser Gly Cys Leu Phe Ser 11e Leu Phe  $260\,$ 8

Pro Leu Phe Ile Ile Ser Ala Asn Glu Ala Lys Thr Pro Gly Lys Ala 275

Ser Asn Leu Val Val Phe Leu 300 Leu Phe Gln Leu Arg Leu Phe Ser 290 ž

Arg Leu Phe His Lys Thr Val Tyr Leu Gln Ser Ala Leu Ser Ser Ser 305

Thr Ser Ala Glu Lys Phe Pro Ser Pro His Pro Ser Pro Ala Lys Leu 335

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Lys Ala Thr Ala Gly His 340 2

(2) INFORMATION FOR SEQ ID NO: 638;

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(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 529 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear

SEQUENCE DESCRIPTION: SEQ ID NO: 638

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Met Ala Lys Phe Met Thr Pro Val Ile Gln Asp Asn Pro Ser Gly Trp 1 5 10 15

Gly Pro Cys Ala Val Pro Glu Glu Gln Phe Arg Asp Met Pro Tyr Gln Pro 20 2

Gly Ala

Ser Ser Gln Phe Gly 60 Lys Val Ala Asp Trp Thr 45 Thr Asn Lys Tyr Phe Ser Lys Gly Asp Arg Leu Gly 35 Thr Tyr Gln Asp Lys Arg Tyr 50 35

Phe 80 Gin Leu Val Asp Thr Ala Arg Thr Gin Lys Thr Ala Tyr Gin Arg Ser Gly Gly Ser Gln Tyr Ala Tyr Phe His Glu Glu Asp Glu 65 6

Agn

Arg Arg Arg Arg Asp Lys Asp 110 Arg Met Arg Phe Ala Gln Arg Asn Leu 100 45

Asn Met Leu Gln Phe Asn Leu Gln Ile Leu Pro Lys Ser Ala Lys Gln 115 S

Gln Lys Lys Phe Gln Lys Gln Phe 140 Ser Gln Lys Pro Arg Asp Ser 155 Trp Asp Gln Lys 150 Le. ile Arg 7 Glu Arg Glu Arg 130 Val Arg Gln Lys Lys 55

Pro Gln Leu Met Lys Met Arg Tyr Leu Glu Val Ser Glu Pro Gln Asp

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Val Lys Glu Glu Met Asp 170

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Arg Ser Asp '

Glu Val

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Arg Gly Ala Val Ile Ala Thr Glu Leu Lys Asn Asn Ser Tyr Lys Leu 420 425 Asn Gly Glu Val 385 Ile Glu Cys Cys Gly Ala Leu Glu Tyr Tyr Asp Lys Ala Phe Asp Arg 195 200 205 Phe Pro Asn Pro Asn Pro Phe Val Glu Asp Asp Met Asp Lys Asn Glu 340 345 His Asn.Phe Ser Gln Gln Cys Leu Arg Met Gly Lys Glu Arg Tyr Asn 325 330 Leu Thr Val Ser Glu Thr Ala Asn Glu Pro Pro Gln Asp Glu Gly Asn 290 295 300 Gly Ser Lys Leu Phe Phe Asp Lys Arg Asp Asn Ser Asp Phe Asp Leu 275 280 285 Ser Cys Thr Arg Ser Val Tyr Ser Trp Asp Ile Val Val Gln Arg Val 260 265 His Thr Val Thr Thr 225 Ile Thr Thr Arg Ser Glu Lys Pro Leu Arg Xaa Xaa Lys Arg Ile Phe 210 215 220 Ile Leu Gly Thr Gln Gln Phe Lys Pro Asn Glu Phe Ala Ser Gln Ile 465 470 475 Ala Arg Trp Thr Cys Cys Ala Leu Leu Ala Gly Ser Glu Tyr Leu Lys 435 440 445 Ile Asp Leu Ile Val Arg Cys Glu His Asp Gly Val Met Thr Gly Ala 370 380 Ile Ala Ser Val Ala Tyr Arg Tyr Arg Ser Gly Lys Leu Gly Asp Asp 355 360 365 Thr Gln Gly Asn Val Phe Ala Thr Asp Ala Ile Leu Ala Thr Leu Met
245 250 Ser Arg His Cys Asn Gly Val Asp Trp Arg Gln Lys Leu Asp Ser Gln
415 Ser Phe Asn Ser Pro Arg Asn Leu Ala Met Glu Ala Thr Tyr Ile Asn 305 310 310 180 Ser Phe Ile Asn Ile Lys Thr Leu Asn Glu Trp Asp 390 395 400 Thr Asp Asp Pro Val Ile Arg Lys Leu Ala Lys 230 235 185

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Ile Cys Met Lys Leu Glu Glu Gly Lys Tyr Leu Ile Leu Lys Asp Pro

Asn Leu Ser Val Glu Asn Ala Trp Gly Ile Leu Arg Cys Val Ile Asp 485 , 490 495 8

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\$ S 50 8 ઝ 30 25 20 15 5 S Val Ala . 145 His Phe Xaa Arg Val Gln Phe His Leu Lys Asn Phe Asp Met Val Ile 115 120 125 Val Tyx Lys Asp Tyx Sex Lys Lys Val Thr Met Ile Asn Ala Ile Pro 130 Ser Lys Tyr Thr Glu Gly Val Gln Ser Leu 165 Thr Glu Trp Pro Pro Phe Val Val Thr Leu Asp Glu Val Glu Leu Ile 100 110 Arg Ser Thr Cys Leu Leu Gln Pro Thr Ser Ser Ala Leu Val Asn Ala 85 90 95 Glu Phe Glu Val Pro Phe Arg Asp Leu Gly Phe Asn Gly Ala Pro Tyr 65 70 75 Phe Lys Asn Phe Ile Glu Lys Val Glu Ala Leu Thr Lys Glu Glu Leu  $50 \ \ 50$ Thr Thr Asp Leu Gly Lys His Gln His Met His Asp Arg Asp Asp Leu 20 25 Lys Lys Arg His Thr Asp Val Gln Phe Tyr Thr Glu Val Gly Glu Ile
1 5 10 15 Asn Lys Gln Val Ile Arg Val Tyr Ser Leu Pro Asp Gly Thr Phe Ser 515 520 520 Thr Ile Val (2) INFORMATION FOR SEQ ID NO: 639: <u>£</u> (i) SEQUENCE CHARACTERISTICS: Ser Leu Asp Asp Pro Glu Gly Phe Phe Glu Gln Gly Gly Trp Ser 180 185 SEQUENCE DESCRIPTION: SEQ ID NO: 639: 500 (A) LENGTH: 194 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear dsy Pro Ile Lys Glu Trp Leu Asn Ser Cys Asp 150 155 505 Asn Trp Thr Lys Ile Met 170 510 160

(2) INFORMATION FOR SEQ ID NO: 640:

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(A) LENGTH: 70 amino acids SEQUENCE CHARACTERISTICS: 3

(B) TYPE: amino acid (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 640:

Arg Ser Gly Leu Gly Leu Gly Ile Thr Ile Ala Phe Leu Ala Thr Leu

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Ile Thr Gln Phe Leu Val Tyr Asn Gly Val Tyr Gln Tyr Thr Ser Pro 20 30

Asp Phe Leu Tyr 11e Arg Ser Trp Leu Pro Cys 11e Phe Ser Gly 45 Gly Val Thr Val Gly Asn Ile Gly Arg Gln Leu Ala Met Gly Val Pro 50 60

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Glu Lys Pro His Ser Asp 65 70

(2) INFORMATION FOR SEQ ID NO: 641: 25

SEQUENCE CHARACTERISTICS

(A) LENGTH: 101 amino acids (B) TYPE: amino acid

(D) TOPOLOGY: linear

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 641:

Val Thr Gin Pro Lys His Leu Ser Ala Ser Het Gly Gly Ser Val Glu 1 5

35

Ile Pro Phe Ser Phe Tyr Tyr Pro Trp Glu Leu Ala Xaa Xaa Pro Xaa 25 30

Val Arg Ile Ser Trp Arg Arg Gly His Phe His Gly Gln Ser Phe Tyr 45 Ser Thr Arg Pro Pro Ser 11e His Lys Asp Tyr Val Asn Arg Leu Phe 50 60 <del>5</del>

Leu Asn Try Thr Glu Gly Gln Glu Ser Gly Phe Leu Arg Ile Ser Asn 65 70 45

Leu Arg Lys Glu Asp Gln Ser Val Tyr Phe Cys Arg Val Glu Leu Asp 85

Thr Arg Arg Ser Gly 100

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(2) INFORMATION FOR SEQ ID NO: 642:

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(1) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 233 amino acids
(B) TYPE: amino acid

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(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 642: (D) TOPOLOGY: linear

Met Glu Ala Gln Gln Val Asn Glu Ala Glu Ser Ala Arg Glu Gln Leu

Gln Xaa Leu His Asp Gln Ile Ala Gly Gln Lys Ala Ser Lys Gln Glu 20

Leu Glu Thr Glu Leu Glu Arg Leu Lys Gln Glu Phe His Tyr Ile Glu 18  $45\ \rm 45$ 2

Glu Asp Leu Tyr Arg Thr Lys Asn Thr Leu Gln Ser Arg Ile Lys Asp 50

Arg Asp Glu Glu Ile Gln Lys Leu Arg Asn Gln Leu Thr Asn Lys Thr 65

2

Leu Ser Asn Ser Gin Ser Glu Leu Glu Asn Arg Leu His Gin Leu 95

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Thr Glu Thr Leu Ile Gln Lys Gln Thr Met Leu Glu Ser Leu Ser Thr 100

Glu Lys Asn Ser Leu Val Phe Gln Leu Glu Arg Leu Glu Gln Gln Met 115 25

Asn Ser Ala Ser Gly Ser Ser Asn Gly Ser Ser Ile Asn Met Ser 110 8

Gly Ile Asp Asn Gly Glu Gly Thr Arg Leu Arg Asn Val Pro Val Leu 145 Phe Asn Asp Thư Glu Thư Asn Leu Ala Gly Met Tyr Gly Lys Val Arg 170

35

Lys Ala Ala Ser Ser Ile Asp Gln Phe Ser Ile Arg Leu Gly Ile Phe 180 6

Leu Arg Arg Tyr Pro Ile Ala Arg Val Phe Val Ile Ile Tyr Met Ala 195 Leu Leu His Leu Trp Val Met Ile Val Leu Leu Thr Tyr Thr Pro Glu 210

Met His His Asp Gln Pro Tyr Gly Lys 225

45

(2) INFORMATION FOR SEQ ID NO: 643

S

(A) LENGTH: 43 amino acids (B) TYPE: amino acid (D) TOPOLOGY: linear (1) SEQUENCE CHARACTERISTICS:

55

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 643

Ile Arg His Glu Gln His Pro Asn Phe Ser Leu Glu Met His Ser Lys  $_{\rm 1}$ 8

S

5

(2) INFORMATION FOR SEQ ID NO: 644:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 63 amino acids

(B) TYPE: amino acid

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 644:

5

Arg His Pro Trp Val Ala Gly Ala Leu Val Gly Val Ser Gly Gly Leu 20Ser Phe Phe Ile Ser Glu Glu Lys Gly His Leu Leu Leu Gln Ala Glu 1 1 15

20

Thr Leu Thr Thr Cys Ser Gly Pro Thr Glu Lys Pro Ala Thr Lys Asn 35 40 45

25

Tyr Phe Leu Lys Arg Leu Leu Gln Glu Met His Ile Arg Ala Asn  $50\,$ 

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PCT/US98/04493

#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

											, 
Auborizzd officer Susan White PCT International Division	This sheet was received with the international application	For receiving Office use only	E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")		D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)	C. ADDITIONAL INDICATIONS (team blank if not applicable)	Date of deposit February 26, 1997	Address of depositary institution (including postal code and country) [1230] Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page $116$ , line N/A
Authorizzd efficer	This sheet was received by the International Bureau on:	For International Bureau use only	resu later (specify the general nature of the indications, e.g., "Accession	·	ARE MADE (If the Indications are not for all designated States)	This information is continued on an additional sheet	Accession Number 97897		tion	Further deposits are identified on an additional sheet	o in the description

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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page $116$	ed to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	lection
Address of depositary institution (including postal code and country)	. (شا
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209043
C. ADDITIONAL INDICATIONS (term blank if not applicable)	(e) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATION	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (now blank If not applicable)	blank if not applicable)
The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Deposit")	lureau later (specify the general nature of the indications, e.g., "Accession
For receiving Office use only	For International Bureau use only
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Authorized officer Susan White PCT International Division	Authorized officer
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PCT/US98/04493

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 119 . line N/A	red to in the description A
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	ilection
Address of depositary institution (including postal code and country)	(A)
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit September 4, 1997	Accession Number 209235
C. ADDITIONAL INDICATIONS (name blank if not applicable)	ble) This information is continued on an additional shect
D. DESIGNATED STATES FOR WHICH INDICATIO	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (name blank if not applicable)	blank if not applicable)
The indications listed below will be submitted to the International Aumber of Deposit?	The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Deposit")
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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

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blank (I not applicable) Sureau later (specify the general nature of the indications, e.g., "Accession	E. SEPARATE FURNISHING OF INDICATIONS (terwe blank (f not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")
NS ARE MADE (If the indications are not for all designated States)	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)
le) This information is continued on an additional sheet	C. ADDITIONAL INDICATIONS (team blank if not applicable)
Accession Number 97898	Date of deposit February 26, 1997
3)	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America
lection	Name of depositary institution American Type Culture Collection
Further deposits are identified on an additional sheet 🔲	B. IDENTIFICATION OF DEPOSIT
ed to in the description	A. The indications made below relate to the microorganism referred to in the description on page 122 , line N/A

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(PCT Rule 13bis)

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)  The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")  For receiving Office use onlyFor International Bureau use only	C. ADDITIONAL INDICATIONS (team blank if not applicable) This information is continued on an additional sheet.	Address of depositary institution (including postal code and country)  12301 Parklawn Drive Rockville, Maryland 20852 United States of America  Date of deposit May 15, 1997  Accession Number 209044	Name of depositary institution American Type Culture Collection	A. The indications made below relate to the microorganism referred to in the description on page 122 , line N/A  B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet
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Susan White PCT International Division

Authorized officer

#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page $126$ , line N/A	red to in the description	A. The indications mad on page 126
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet	B. IDENTIFICATIO
Name of depositary institution American Type Culture Collection		Name of depositary insti
Address of depositary institution (including postal code and country)	(y)	Address of depositary in
12301 Parklawn Drive Rockville, Maryland 20852 United States of America		12301 Parklawn Drivo Rockville, Maryland United States of Amer
Date of deposit February 26, 1997	Accession Number 97899	Date of deposit May
C. ADDITIONAL INDICATIONS (new blant if not applicable)	it.) This information is continued on an additional sheet	C. ADDITIONAL IN
ESIGNATED STATES FOR WHICH INDICATIO	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the Indicators are not for all designated States)	D. DESIGNATED ST
E. SEPARATE FURNISHING OF INDICATIONS (sear blank if not applicable)	blank if not applicable)	E. SEPARATE FURN
dications listed below will be submitted to the International or of Depart?	The indications listed below will be submitted to the international Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	The indications listed belo Number of Deposit?
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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

on page 126 , line N/A	Ted to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	illection
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	(17)
Date of deposit May 15, 1997	Accession Number 209045
C. ADDITIONAL INDICATIONS (new blank if not applicable)	(bb) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIO	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are most for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (team blank () not applicable) The indications listed below will be submitted to the International Bureau later (specify the Number of Deposit")	E. SEPARATE FURNISHING OF INDICATIONS (acre blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")
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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

Authorized officer Supen White PCT International Division	This sheet was received with the international application	For receiving Office use only	E. SEPARATE FURNISHING OF INDICATIONS (term blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (have blank if not applicable)	Date of deposit April 28, 1997 Ac	12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Address of depositary institution (including postal code and country)	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page 130 , line N/A
Authorized officer	This sheet was received by the International Bureau on:	For International Bureau use only	th (I not applicable)  au later (specify the general nature of the indications, e.g., "Accession	ARE MADE (If the indications are not for all designated States)	This information is continued on an additional sheet	Accession Number 209011			on	Further deposits are identified on an additional sheet	in the description

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(PCT Rule 13bis)

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

Authorizze officer Sugan White PCT International Division	FOR FECEIVING OFFICE USE ONLY  This sheet was received with the international application	Englishing Office we only	E. SEPARATE FURNISHING OF INDICATIONS (traw blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (leave blank if not applicable)	Date of deposit February 26, 1997	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page $131$ , line N/A
Authorized officer	This sheet was received by the International Bureau on:	For Intermetional Burson use only	lank (I not applicable) ureau later (specify the general nature of the indications, e.g., "Accession	S ARE MADE (I) the indications are not for all designated States)	) This information is continued on an additional sheet	Accession Number 97900		ction	Funher deposits are identified on an additional sheet 📋	to in the description

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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page $137$ , line $N/A$	A. The	A. The indications on page 131
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet	B. IDE	B. IDENTIFICA'
Name of depositary institution American Type Culture Collection	Name o	Name of depositary
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Address 12301 Rockvi United	Address of deposita 12301 Parklawn E Rockville, Maryla United States of A
Date of deposit February 26, 1997 Accession Number 97901	Date of	Date of deposit N
C. ADDITIONAL INDICATIONS steam blank if not applicable) This information is continued on an additional sheet	C. AD	C. ADDITIONA
	.	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	D. DES	D. DESIGNATEI
E. SEPARATE PURNISHING OF INDICATIONS	d as	E SEPARATE EI
The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Deposit")	The indicate the indicate of t	The indications listed Number of Deposit")
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### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page 131     , line N/A	cd to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	llection
Address of depositary institution (including postal code and country)	(6)
12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit May 15, 1997	Accession Number 209046
C. ADDITIONAL INDICATIONS (teave blank if not applicable)	16) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIO	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)
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E. SEPARATE FURNISHING OF INDICATIONS (naw blank I not applicable)	blank if not applicable)
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., Number of Deposit")	lurcau later (specify the general nature of the indications, e.g., "Accession
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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

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DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	D. DESIGNATED STATES FOR WHICH INDICATION
(e) This information is continued on an additional sheet 🔲	C. ADDITIONAL INDICATIONS (teame blank if not applicable)
Accession Number 209047	Date of deposit May 15, 1997
3	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America
lection	Name of depositary institution American Type Culture Collection
	B. IDENTIFICATION OF DEPOSIT
ed to in the description	A. The indications made below relate to the microorganism referred to in the description on page 137 , line N/A

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<b>,</b>	
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### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

Authorized officer Sugan White	This sheet was received with the international application	For receiving Office use only	E. SEPARATE FURNISHING OF INDICATIONS (near blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify it number of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)	C. ADDITIONAL INDICATIONS (leave blank if not applicable)	Date of deposit May 22, 1997	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page $-137$ , line $-\rm N/A$
Authorized officer	This sheet was received by the international Bureau on:	For International Bureau use only	B. SEPARATE FURNISHING OF INDICATIONS (serve blank () not applicable) The indications listed below will be submitted to the International Bureau later (speelly the general nature of the Indications, e.g., "Accession Number of Depart")	IS ARE MADE (If the indications are not for all designated States)	e) This information is continued on an additional sheet 🔲	Accession Number 209076		etion	Further deposits are identified on an additional sheet	to in the description

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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

	(PCT Rule 1361s)		
A. The indications made below relate to the microorganism referred to in the description on page $\frac{140}{}$ , line N/A	rrred to in the description A		A. The indicatio on page
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet.		B. IDENTIFIC
Name of depositary institution American Type Culture Collection	ollection		Name of deposita
Address of depositary institution (including postal code and country)	nty)		Address of depos
12301 Parklawn Drive Rockville, Maryland 20852 United States of America			12301 Parklawi Rockville, Mari United States o
Date of deposit August 21, 1997	Accession Number 209215		Date of deposit
C. ADDITIONAL INDICATIONS from blank if not applicable	able) This information is continued on an additional sheet		C. ADDITION
		•	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	NS ARE MADE (if the indications are not for all designated States)		D. DESIGNAT
÷			
E. SEPARATE FURNISHING OF INDICATIONS (neare blank if not applicable)	blank if not applicable)		E. SEPARATE
The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Deposit")	Bureau later (specify the general nature of the indications, e.g., "Accession		The indications list Number of Deposit')
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## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

A. The indications made below relate to the microorganism referred to in the description on page $160$ , line $N/A$	referred to in the description N/A
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	ollection
Address of depositary institution (including postal code and country) 1230! Parklawn Drive	uiry)
Rockville, Maryland 20852 United States of America	
Dale of deposit February 26, 1997	Accession Number 97904
C. ADDITIONAL INDICATIONS (team blank if not applicable)	able) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATION	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (nowe blank if not applicable)	e blank if not applicable)
The indications listed below will be submitted to the International Number of Deposit?	The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Deposit")
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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

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DESIGNALED STALES FOR WHICH INDICATIONS AND MADE (9 instructions are not for an actignates states)	D. DESIGNALED STATES FOR WHICH INDICATION
le) This information is continued on an additional sheet	C. ADDITIONAL INDICATIONS (leave blank if not applicable)
Accession Number 209139	Date of deposit July 3, 1997
(4)	Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America
lection	Name of depositary institution . American Type Culture Collection
Further deposits are identified on an additional sheet	B. IDENTIFICATION OF DEPOSIT
ed to in the description	A. The indications made below relate to the microorganism referred to in the description on page 154 , line N/A

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(PCT Rule 13bis)

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

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E. SEPARATE FURNISHING OF INDICATIONS (term blank (f not applicable) The indications listed below will be submitted to the International Bureau later (upscify it Number of Deposit')	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (1) the indications are not for all designated States)		C. ADDITIONAL INDICATIONS (leave blank if not applicable)	Date of deposit May 15, 1997	(1230) Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution  American Type Culture Collection	, IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page 153 , line N/A
E. SEPARATE FURNISHING OF INDICATIONS (seaw blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the Indications, e.g., "Accession Number of Departs")	S ARE MADE (If the indications are not for all designated States)		e) This information is continued on an additional sheet	Accession Number 209049		ection	Further deposits are identified on an additional sheet	d to in the description

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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page $153$ , ine N/A	rred to in the description A
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	ollection
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	ury)
Date of deposit February 26, 1997	Accession Number 97903
C. ADDITIONAL INDICATIONS (leave blank if not applicable)	ible) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATION	DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (now blank if not applicable)	blank if not applicable)
The indications listed below will be submitted to the International Number of Deposit")	The indications listed below will be submitted to the International Bureau later (specify the general nature of the indication, e.g., "Accession Number of Deposit")
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## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

A. The indications made below relate to the microorganism referred to in the description on page $142$ , line N/A	I to in the description
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet
Name of depositary institution American Type Culture Collection	ction
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	
Date of deposit June 12, 1997	Accession Number 209119
C. ADDITIONAL INDICATIONS (teaw blank if not applicable)	) This information is continued on an additional sheet
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the Indications are not for all designated States)	S ARE MADE (if the Indications are not for all designated States)
E. SEPARATE FURNISHING OF INDICATIONS (new blank if not applicable)	lank if not applicable)
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	ureau later (specify the general nature of the indications, e.g., "Accession
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#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

Authorized officer Sussan Willie PCT International Division	For receiving Office use only  This sheet was received with the international application	E. SEPARATE FURNISHING OF INDICATIONS (team blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications, e.g., "Accession Number of Deposit")	D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (If the indications are not for all designated States)	C. ADDITIONAL INDICATIONS (leave blank if not applicable)	Date of deposit February 26, 1997 A	Address of depositary institution ( <i>including postal code and country</i> ) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	Name of depositary institution American Type Culture Collection	B. IDENTIFICATION OF DEPOSIT	A. The indications made below relate to the microorganism referred to in the description on page 146 , line N/A
Authorized officer	For International Bureau use only  This sheet was received by the International Bureau on:	ank (frot applicable) ireau later (specify the general nature of the indications, e.g., "Accession	ARE MADE (If the indications are not for all designated States)	This information is continued on an additional since.	11		ion	Further deposits are identified on an additional sheet	) in the description

## INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

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Authorizzd officer	This sheet was received by the International Bureau on:	For International Bureau use only	nk (I not applicable)	ARE MADE (If the Indications are not for all designated States)	This information is continued on an additional sheet.			on	Further deposits are identified on an additional sheet	in the description

#### INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

Piene of depositary institution American Type Culture Collection Address of depositary institution Address of depositary institution Address of depositary institution Address of depositary institution Address of depositary institution (including pootal code and country) L200 Packlawn Drive Reckville, Mary 15, 1997  Date of deposit May 15, 1997  C. ADDITIONAL INDICATIONS (now bend fram applicable)  D. DESIGNATED STATES FOR WHICH INDICATIONS (now bend fram applicable)  E. SEPARATE FURNISHING OF INDICATIONS (now bend fram applicable) This indications listed below will be submitted to the International Bureau later (pacify for general feature of the indications at the Address of the Address of the Address of the International Bureau later (pacify for general feature only address of the International Bureau later (pacify for general feature only address of the International Bureau and conly address of the International Bureau and conly address of the International Bureau and conly address of the International Bureau and conly address of the International Bureau and conly address of the International Bureau and conly address of the International Bureau and conly address of the International Bureau and the International Bureau and the International Bureau and Conly Authorited officer  Suggest Whithe PCT International Daddion  Authorited officer  Suggest Whithe	A. The indications made below relate to the microorganism referred to in the description on page 160     Inc N/A	ed to in the description
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# INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

A. The indications made below relate to the microorganism referred to in the description on page $\sim 142$ , line N/A	eferred to in the description N/A
B. IDENTIFICATION OF DEPOSIT	Further deposits are identified on an additional sheet 🛛
Name of depositary institution American Type Culture Collection	ollection
Address of depositary institution (including postal code and country) 12301 Parklawn Drive Rockville, Maryland 20852 United States of America	н(гу)
Date of deposit February 12, 1998	Accession Number 209627
C. ADDITIONAL INDICATIONS (teaw blank if not applicable)	able) This information is continued on an additional sheet
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Form PCT/RC/1114 (Italy 1992)	

#### What Is Claimed Is:

1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:

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- (a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;

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- (c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;
- (d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a
  polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z,
  which is hybridizable to SEQ ID NO:X;

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- (e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, having biological activity;
- (f) a polynucleotide which is a variant of SEQ ID NO:X;

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- (g) a polynucleotide which is an allelic variant of SEQ ID NO:X;
- (h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;
- (i) a polynucleotide capable of hybridizing under stringent conditions to any
- 25 one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.
- The isolated nucleic acid molecule of claim 1, wherein the
   polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.
- 3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

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- 4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.
- 5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.
- The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the Nterminus.

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 A recombinant vector comprising the isolated nucleic acid molecule of claim 1.

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- A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.
- A recombinant host cell produced by the method of claim 8.

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- 10. The recombinant host cell of claim 9 comprising vector sequences.
- 11. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:

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- (a) a polypeptide fragment of SEQ ID NO: Y or the encoded sequence included in ATCC Deposit No:Z;
- (b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z, having biological activity;
- (c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;
- (e) a secreted form of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

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(f) a full length protein of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

- (g) a variant of SEQ ID NO:Y;
- (h) an allelic variant of SEQ ID NO: Y; or
- (i) a species homologue of the SEQ ID NO:Y.
- The isolated polypeptide of claim 11, wherein the secreted form or the
  - in comprises sequential amino acid deletions from either the C-terminus full len or th

'body that binds specifically to the isolated polypeptide of

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\*xpresses the isolated polypeptide of claim

4.

15. A method.

· comprising:

said polypeptide is expressed; and (a) culturing the recon.

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under conditions such that

(b) recovering said polypeptide.

The polypeptide produced by claim 15. 16.

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- comprising administering to a mammalian subject a therapeutically effective amount of A method for preventing, treating, or ameliorating a medical condition, the polypeptide of claim 11 or the polynucleotide of claim 1.
- A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising: . 8

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- (a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and
- (b) diagnosing a pathological condition or a susceptibility to a pathological
  - condition based on the presence or absence of said mutation. 3
- 19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

(a) determining the presence or amount of expression of the polypeptide of

claim 11 in a biological sample; and

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(b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

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A method for identifying a binding partner to the polypeptide of claim 11

comprising:

(a) contacting the polypeptide of claim 11 with a binding partner; and

(b) determining whether the binding partner effects an activity of the

polypeptide.

The gene corresponding to the cDNA sequence of SEQ ID NO: Y.

A method of identifying an activity in a biological assay, wherein the

method comprises:

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(a) expressing SEQ ID NO:X in a cell;

(b) isolating the supernatant;

(c) detecting an activity in a biological assay; and

(d) identifying the protein in the supernatant having the activity.

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The product produced by the method of claim 22. 23.